

1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the
Genbank flat file format but are available as part
of this entry's ASN.1 file.

Double stranded (DS) coverage: 75.5%
DS or two chemistry coverage: 98.9%
Single stranded regions: 3

Sequence Validation:
This sequence has been validated by Multiple Complete Digest
Mapping. Comparison of the experimentally derived map digest
fragments with sequence-predicted fragments is given below.
Small fragments below a variable cutoff (approximately 400-600bp)
are not mapped and hence do not appear in the table. There are no
significant remaining discrepancies between the experimental and
predicted values. Uniquely ordered fragment groups are separated
by dashed lines.

Map	Seq	Map	Seq	Map	Seq
BglII		HindIII		MsiI	
1069.11	1050.00	889.55	866.00	30541.40	30653.00
20320.67	20855.00	1050.18	1015.00	3379.08	3231.00
2171.50	2147.00	7268.78	7196.00		
2560.20	2531.00	10085.80	9992.00		
4335.42	4269.00	11212.78	11131.00		
2698.62	2628.00				
1927.50	1887.00				
3130.46	3090.00				
2166.69	2129.00				
2044.57	2005.00				

FEATURES
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/db_xref="taxon:9606"
/chromosome="6"
/map="6p21"
/sub_clone="UMGC:370M23.002"
/clone_lib="Research Genetics BAC Library"
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complement(4999..5277)
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6285..6572
/rpt_family="Alu"
complement(6972..7050)
/rpt_family="MLT1"
7286..7584
/rpt_family="Alu"
complement(8164..8609)
/rpt_family="Alu"
complement(21287..21895)
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22715..22957
/rpt_family="Alu"
25510..25802
/rpt_family="Alu"
27835..28010
/rpt_family="MER20"
31295..31594
/rpt_family="Alu"
33515..33767
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repeat_region 37372..37648
/rpt_family="Alu"
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/rpt_family="MER3"
repeat_region 39583..40010
/rpt_family="Alu"
repeat_region 40046..40156
/rpt_family="Alu"
repeat_region 43194..43372
/rpt_family="MER5"
variation 43325
/note="clonal variation with 3' overlapping clone"
variation 44149
/note="clonal variation with 3' overlapping clone"
variation 44491
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variation 45760
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variation 45900
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variation 46851
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variation 47032
/note="clonal variation with 3' overlapping clone"
variation 47240..47256
/note="clonal variation with 3' overlapping clone -
insertion of 17bp repeat"

BASE COUNT 11556 a 11489 c 12284 g 11994 t
ORIGIN

Query Match 39.3%; Score 92; DB 31; Length 47323;
Best Local Similarity 100.0%; Pred. No. 1.14e-47;
Matches 92; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 35465 CAGGCGCGGTATGACTTGCACCTGACGACGCTTTTGCACAAATCTCCT 35524
QY 109 CAGGCGCGGTATGACTTGCACCTGACGACGCTTTTGCACAAATCTCCT 168
Db 35525 ATGAGTCACCTCTCGAATTGCTGAAAG 35556
QY 169 ATGAGTCACCTCTCGAATTGCTGAAAG 200

RESULT 2
LOCUS 166494 7218 bp DNA PAT 23-DEC-1997
DEFINITION Sequence 14 from patent US 5670367.
ACCESSION 166494
NID 92724471
VERSION 166494.1 GI:2724471
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 7218)
AUTHORS Dörner, F., Schefflinger, F. and Falkner, F. Gunter.
TITLE Recombinant fowlpox virus
JOURNAL Patent: US 5670367-A 14 23-SEP-1997;
FEATURES
source
1..7218
/organism="unknown"
BASE COUNT 1944 a 1491 c 1486 g 1929 t 368 others
ORIGIN

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Query Match      20.5%; Score 48; DB 25; Length 7218;
Best Local Similarity 1.8%; Pred. No. 1,55e-15;
Matches 4; Conservative 130; Mismatches 86; Indels 0; Gaps 0;

Db 1221 YYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYY 1280
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Qy 63 CTGCCCTTCTCTCTCTCTCTGACCTCTGACCTCATGACAGGCGCGGTATG 122
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1281 YYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYY 1340
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Qy 123 ACTTGCACTGACGTGACGAGCTTTTCTGACAAATCTCTCTATGAGTCCAGCTTC 182
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1341 YYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYY 1400
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Qy 183 CTGGAATTCCTGAAAGCTCTGCCCTCCCTCCATCTCCCTAGGACACAGCTCC 242
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1401 YYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYY 1440
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Qy 243 CTCACACATGCAAGATCTCACACACCATGTTGTCCTACACA 282
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 3
LOCUS AC005369 74371 bp DNA PRI 01-AUG-1998
DEFINITION Homo sapiens chromosome 5, BAC clone 119j3 (LBNL H175), complete
sequence.
ACCESSION AC005369
NID 93367505
VERSION AC005369.1 GI:3367505
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
          Primates; Catarrhini; Hominoidea; Homo.
REFERENCE 1 (bases 1 to 74371)
AUTHORS Kimerly,W., Bondoc,M., Cheng,J., Connolly,K.S., Gunning,K.M.,
          Kader,K., Kader,K., Miguel,T., Pfluck,S., Pollard,M.,
          Rojeski,H., Subramanian,S. and Martin,C.H.
          Submitted (01-AUG-1998) Human Genome Center, DOE Joint Genome
          Institute, Lawrence Berkeley National Laboratory, MS 74-157,
          Berkeley, CA 94720, U.S.A.
          Sequence submitted by:
          DOE Joint Genome Institute.
          Location/Qualifiers
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              1..74371
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
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                /clone="119j3"
                /chromosome="5"
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            repeat_region
              893..1030
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            repeat_region
              2295..2438
                /rpt_family="Alu"
            repeat_region
              2818..2859
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            rpt_unit=GT
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            /standard_name="RF"
            /note="65% & 69% protein identity GenPept:U22377"

repeat_region 3431..3724
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repeat_region 5327..5602
            /rpt_family="Alu"
repeat_region 6586..6956
            /rpt_family="L1"
repeat_region 6647..6684
            /note="(CA)19"
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            /rpt_unit=CA
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            /rpt_unit=AC
            /rpt_family="Alu"
repeat_region 8258..8503
            /rpt_family="Alu"
            /rpt_type=tandem
            /rpt_unit=AC
            /rpt_family="Alu"
repeat_region 9070..9387
            /rpt_family="Alu"
            /rpt_type=tandem
            /rpt_unit=AC
            /rpt_family="Alu"
repeat_region 9740..9845
            /rpt_family="MER42"
            /rpt_type=tandem
            /rpt_unit=AC
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repeat_region 10440..11015
            /rpt_family="Alu"
            /rpt_type=tandem
            /rpt_unit=AC
            /rpt_family="Alu"
repeat_region 11950..12250
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            /rpt_family="Alu"
repeat_region 12057..12085
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            /rpt_unit=AA
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repeat_region 12365..12645
            /rpt_family="Alu"
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repeat_region 13727..13750
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repeat_region 13783..14024
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repeat_region 14175..14470
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            /rpt_type=tandem
            /rpt_unit=AC
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            /rpt_family="Alu"
repeat_region 16993..17085
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            /rpt_type=tandem
            /rpt_unit=AC
            /rpt_family="Alu"
repeat_region 17678..18276
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            /rpt_unit=AC
            /rpt_family="Alu"
repeat_region 21736..22035
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            /rpt_type=tandem
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Note: remainder of annotations omitted.

misc_feature
    /note="78%-100% protein identity GenPept:U01933/"
    complement(38069..38215)
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Query Match      14.1% Score 33; DB 31; Length 74321;
Best Local Similarity 17.9%; Pred. No. 9.92e-06;
Matches 19; Conservative 53; Mismatches 33; Indels 1; Gaps 1;

Db 15901 TTTTTCGASKSRGWSCKCCKCTSTSCSOMSRKSRRMGVGYNSMKTRCAAMWTKSSKC 15960
|||||::: :: :: : |::: ::: ::: ::: ::: ::: ::: ::: ::: ::: :::
Oy 149 TTTTTCGACAAATTCCTCATAGCACCGCTTCGGATTGCATAAGCTCTGCCCT 208
|||||::: ::: ::: ::: ::: ::: ::: ::: ::: ::: ::: ::: ::: ::: :::
Db 15961 WWSYSTRMMKYCSGYCYSGSKRGWKRCNSMYWTGYSYRYWMS 16006
||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||:
Oy 209 CCTCTCCATCTCCTTCAGGACCA-GCGTCACCCTCCACCAATGC 253
||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||: ||:

RESULT 4
LOCUS HUAC004787 216021 bp DNA PRI 24-JUN-1998
DEFINITION Homo sapiens chromosome 16 BAC clone C19875K-A-952F10, complete sequence.
ACCESSION AC004787
IID 9337381
VERSION AC004787.1 GI:3337381
SOURCE HGS.
ORGANISM human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (bases 1 to 216021)
Adams,M.D., Loftus,B.J., Zhou,L., Crosby,M., Fuhrmann,J., Mason,T.M., Brandon,R., Kim,U.Y., Kevlavage,A.-R. and Venter,J.C.
Homo sapiens Chromosome 16 BAC clone C19875K-A-952F10
Unpublished
2 (bases 1 to 216021)
Adams,M.D. and Loftus,B.J.
Direct Submission
Submitted (02-JUN-1998) The Institute for Genomic Research, 9712 Medical Center Dr, Rockville, MD 20850, USA, Email: b.loftus@tigr.org
3 (bases 1 to 216021)
Adams,M.D. and Loftus,B.J.
Direct Submission
Submitted (24-JUL-1998) The Institute for Genomic Research, 9712 Medical Center Dr., Rockville, MD 20850, USA
On Jul 24, 1998 this sequence version replaced gi:3241936.
Addresses all correspondence to: Mark Adams The institute for Genomic Research 9712 Medical Center Dr. Rockville, MD 20850, USA e-mail address: humgen@tigr.org. The orientation of the sequence is from Sfp end to 3' end. Genes were identified by a combination of five methods including: XGRail (available by anonymous ftp from arthur.spm.ornl.gov), Genefinder (Phil Green, University of Washington), Genscan (Chris Burge, htp://gnome.stanford.edu/~chris/GENSCAN.html), searches of the complete sequence against a peptide database, and the Human gene Index database at NIG (http://www.tigr.org/tdb/hgi.html).
Genes without peptide homology having spliced EST hits are termed 'unknown gene product'. Genes encoding tRNAs are predicted by tRNAscan-SE (Sean Eddy, http://genome.wustl.edu/eddy/tRNAscan-SE/) Location/Variants
1..216021
/organism="Homo sapiens"
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/chromosome="16"
/map="#16g21.22"
/cclone="A-952F10"
27765..27872
/note="7766, STSL-CSR1-27g3-uA, CSR1-27g3-uZ, Chr. -, Homo sapiens"
/db_xref="dbSTS:GC02280"
73826..73943

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[illegible]

	SOURCE	Unknown.
	ORGANISM	Unknown.
	REFERENCE	Unclassified. 1 (bases 1 to 965)
	AUTHORS	Wallace,I.,Paul, Harris,W.J., Carr,F.J., Old,L.J., Wels,S. and Kitamura,K.
	TITLE	Recombinant human anti-Lewis b antibodies
	JOURNAL	Patent: US 5795961-A 22-18-AUG-1998;
	FEATURES	Location/Qualifiers source 1..965
	BASE COUNT	/organism="unknown" ORIGIN 192 a 170 c 226 g 205 t 172 others
	Query Match	12.0%: Score 28; DB 25; Length 965; Best Local Similarity 16.3%; Pred.No.9,45e-03; Matches 26; Conservative 65; Mismatches 69; Indels 0; Gaps 0;
Db	771 AARTGGCTKKKTRHTTVVSGGVNSTSTASDYTTSYMGWVRGMDGYGGYTNNKR	830
Cp	225 AAGCGAGATGGAGGAGGAGGACAGACTTTCAGCAATCCAGAGACTGCATCATAAG	166
Db	831 GRTYTAADTSNBSBSVPAADLVAVYCVAGRSDSDGDYWGCTTYSSHTVDMTSSS	890
Cp	165 AGGAATTGTGCAAAAGACGCTTCCTCACCTTGCAATTCACCGGCCCTGTTC	106
Db	891 ASVGDRVTTCRSSSTTHNGNTYYTKGRAKRYVNSRGVS	930
Cp	105 CATGTGACCTGCCAAGAGAGGTCACAAGAGACAAAGGGG	66
RESULT	7	
LOCUS	128278	215 bp DNA PAT 30-OCT-1996
DEFINITION	Sequence 5 from patent US 5569830.	
ACCESSION	128278	
NID	g1819054	
VERSION	128278.1	GI:1819054
KEYWORDS	.	
SOURCE	Unknown.	
ORGANISM	Unknown.	
REFERENCE	1 (bases 1 to 215)	Unclassified.
AUTHORS	Bennett,A.; Labavitch,J.M.; Powell,A. and Stoltz,H.	
TITLE	Plant inhibitors of fungal polygalacturonases and their use to control fungal disease	
JOURNAL	Patent: US 5569830-A 5-29-OCT-1996;	
FEATURES	Location/Qualifiers source 1..215	
BASE COUNT	15 a 8 c 25 g 26 t 141 others	
ORIGIN		
Query Match	11.5%: Score 27; DB 25; Length 215; Best Local Similarity 20.0%; Pred.No.3,51e-02; Matches 22; Conservative 42; Mismatches 45; Indels 1; Gaps 1;	
Db	18 CNDKAKKGNITSSWTDDCNFTGVCDDITTRYRVNDSGHNKISSANTNYGNNVGA	77
Oy	89 CTTTGCAAGCTCATGGACAGCGCGGGTAGACCTTTCGACATCGACAGCTGAA	147
Db	78 THYTTTNVSGADSKVTIDYSYNASGTSNSSNGDTGDNRSAGDSSYSTAM	127
Oy	148 CTTTCTGACAAATTCCTCTATGATGCACGCTCTCTGTGAATCTCTGAA	197
RESULT	8	
LOCUS	HUAC004787.216021 bp DNA PRI 24-JUL-1998	
DEFINITION	Homo sapiens Chromosome 16 BAC clone CIT987SK-A-952F10, complete sequence.	
ACCESSION	AC004787	
NID	93337381	
VERSION	AC004787.1	GI:3337381
KEYWORDS	HTG.	

	SOURCE	human.
	ORGANISM	Homo sapiens
		Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
		Primates; Catarrhini; Hominoidea; Homo.
REFERENCE	TITLE	1 (bases 1 to 216021)
AUTHORS		Adams,M.D., Loftus,B.J., Zhou,L., Crosby,M., Fuhrmann,J.,
		Mason,T.M., Brandon,R., Kim,U.U., Keitelage,A.R. and Venter,J.C.
JOURNAL		Homo sapiens Chromosome 16 BAC clone C1987/SK-952F10
REFERENCE	TITLE	unpublished
AUTHORS		2 (bases 1 to 216021)
JOURNAL		Submitted Submission
REFERENCE	TITLE	Direct Submission
AUTHORS		Submitted (02-JUN-1998) The Institute for Genomic Research, 9712
JOURNAL		Medical Center Dr, Rockville, MD 20850, USA, Email:
COMMENT		bjloftus@tigr.org
		3 (bases 1 to 216021)
		Adams,M.D. and Loftus,B.J.
		Direct Submission
		Submitted (24-JUL-1998) The Institute for Genome Research, 9712
		Medical Center Dr., Rockville, MD 20850, USA
		On Jul 24, 1998 this sequence version replaced gi:3241936.
		Address all correspondence to: Mark Adams The Institute for Genomics
		Research 9712 Medical Center Dr, Rockville, MD 20850, USA e-mail
		address: humgenet@tigr.org. The orientation of the sequence is from
		snp end to 3' end. Genes were identified by a combination of five
		methods including: XSRail (available by anonymous ftp from
		arthur.epm.ornl.gov), GeneFinder (Phil Green, university of
		Washington), Gensean (Chris Bugge,
		http://gnomic.stanford.edu/~chris/GENSEANW.html) searches of the
		complete sequence against a peptide database, and the Human gene
		index database at TIGR (http://www.tigr.org/tdb/hgi.html).
		Genes without peptide homology having spliced EST hits are termed
		'Unknown gene product'. Genes encoding tRNAs are predicted by
		tRNAscan-se (Sean Eddy, http://genome.wustl.edu/eddy/tRNAscan-SE/)
FEATURES	source	location/Organisms
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		/chromosome="16"
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		/clone="A-952F10"
STS		27765.. 27872
		/note="766, STS1-CSRL-2793-UA/CSRL-2793-UZ, Chr. -, Homo
		sapiens"
		/db_xref="dbSTS:G02280"
STS		73826.. 73943
		/note="7608, STS1-CSRL-24g1-UA/CSRL-24g1-UZ, Chr. -, Homo
		sapiens"
		/db_xref="dbSTS:G02122"
STS		175801.. 175945
		/note="16084, CHUC.GCT10B02, Chr. -, Homo sapiens"
		/db_xref="dbSTS:G09703"
STS		175810.. 175945
		/note="16316, CHUC.GCT15C04, Chr. -, Homo sapiens"
		/db_xref="dbSTS:G09935"
STS		199463.. 199572
		/note="9824, WI-3555, Chr. 16, Homo sapiens"
		/db_xref="dbSTS:G04338"
BASE COUNT		60960 a 51778 c 49172 g 55987 t 124 others
ORIGIN		
Query Match		11.5% Score 27; DB 31; Length 216021;
Best Local Similarity		13.8%; Pred.No. 3,51e-02;
Matches		18; Conservative 63; Mismatches 48; Indels 1; Gaps 1;
Dn	1457	KYMMKSMRRARRSGAGKKKKYYCYCYYYYYYTCMGRAAAAATYRRCVAMTY 1516
Cp	227	TGAAGAAGATGGAGAGGACGACACTTTTCAAGCAATTCCAGAAAGCTGCATCAT 168
Dd	1517	YMRARAKTYTAYRRRGCAASYSK-KTMANTYAAATTTCAAAAAAAMTTTTYMMATYRK 1575
Cp	167	GAGGAAATTTGTCTGAAAGAACCTCTCCACTTGCGTTCGAAAGAGTCATACCCTGGCCCTGT 108

RESULT	9	E04076	565 bp	RNA	PAT	29-SEP-1997
LOCUS		gDNA encoding envelope region of type C hepatitis virus.				
DEFINITION		E04076				
ACCESSION		E04076				
NID		92172286				
VERSION		E04076.1 GI:2172286				
KEYWORDS		JP 1992349885-A/1.				
SOURCE		Hepatitis C virus.				
ORGANISM		Hepatitis C virus				
		Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;				
		Hepatitis C-like viruses.				
		1 (bases 1 to 565).				
REFERENCE		Moriyama,T., Chayama,K., Kumada,H. and Ichikawa,Y.				
AUTHORS		NONCODIC ACID FRAGMENT WITHIN ENVELOPE REGION OF HEPATITIS C VIRUS				
TITLE		AND METHOD FOR DETECTING THE SAME				
JOURNAL		Patent: JP 1992349885-A 1 04-DEC-1992;				
COMMENT		TELEJIN LTD				
		OS Hepatitis C virus				
		PN JP 1992349885-A/1				
		PD 04-DEC-1992				
		PF 29-MAY-1991				
		PI MORIYAMA TSUNAE, CHAYAMA KAZUOKI, KUMADA HIROMITSU, PI				
		ICHIKAWA YUTARO				
		PC, CI2N15/10,C12Q1/68,C12Q1/70//C12N15/11:				
		CC strandedness: Single;				
		CC topology: Linear;				
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		/db_xref="taxon:11103"				
BASE COUNT		60 a 93 c 107 g 85 t 220 others				
ORIGIN						
Query Match		10.7% Score 25; DB 25; Length 565;				
Best Local Similarity		34.1%; Pred.No.4.53e-01;				
Matches		30; Conservative 29; Mismatches 28; Indels 1; Gaps 1;				
Db		400 GGRCTGCTTTTCVTRTYTCCABYTGTACTCTTBMBCTMGSGIVAGDACDRY 459				
Oy		54 GGATGTGGCGTCCGCCCTTGCTCCTCCTCTGCACCCTCGTGGCACTCATATGGAACGG 113				
Db		460 CARGTGYAAATGTCCTDMYTAATSCYG 487				
Oy		114 CCGGCTA-TGACTTTGGCAACAGAAGCTG 140				
RESULT	10	AB002302	6252 bp	mRNA	PRI	13-FEB-1999
LOCUS		Human mRNA for KIA00304 gene, complete cds.				
DEFINITION		AB002302				
ACCESSION		92224548				
NID		AB002302.1 GI:2224548				
VERSION		KIA00304.				
KEYWORDS		Homo sapiens male brain cDNA to mRNA, clone_1lb:pbluescriptII SK				
SOURCE		plus clone:H6016.				
ORGANISM		Homo sapiens				
		Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;				
		Primates; Catarrhini; Hominoidea; Homo.				
		1 (bases 1 to 6252)				
REFERENCE		Nagase,T., Ishikawa,K., Seki,N., Nakajima,D., Ohira,M.,				
AUTHORS		Miyajima,N., Kotani,H., Nomura,N. and Ohara,O.				
TITLE		Direct Submission				
JOURNAL		Submitted (28-MAR-1997) to the DDBJ/EMBL/GenBank databases. Nobuo				
		Nomura, Kazusa DNA Research Institute, Gene Structure 1: 1532-3				
		Yana, Kistatazu, Chiba 292, Japan (E-mail:cdnainfo@kazusa.or.jp,				
		URL:http://www.kazusa.or.jp, Tel:+81-438-52-3930,				
		Fax:+81-438-52-3931)				
REFERENCE		2 (sites)				

[illegible]

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misc_feature 570..587
/note="H4 specific box"
promoter 597..604
/note="TATA-box"
precursor_RNA 627..1025
/note="pot. primary transcript"
precursor_RNA 628..1025
/note="pot. alt. primary transcript"
CDS 660..971
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/codon_start=1
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/db_xref="PID:951311"
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/db_xref="MGD:MGI:96099"
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/translation="MSGRGGRGGLGKGGAKRRKRVLRNDNIQITKPAIRRLARRGV
KRISGLIEETRGVLKVFLENVINDAVYTEHAKRKTAMDVYRLKRGRTLYGFG
G"

BASE COUNT 414 a 340 c 284 g 349 t 291 others
ORIGIN

Query Match 9.8% Score 23; DB 32; Length 1678;
Best Local Similarity 78.0%; Pred. No. 5.25e+00;
Matches 32; Conservative 0; Mismatches 9; Indels 0; Gaps 0;
DB 1028 GAATTCTTAAGCTGTGCTGCTCCCTCACCCTTC 1068
||||| ||| ||||| ||| | ||||| |||||
OY 186 GAATTCCTGAAGCTGTGCTGCTCCCTCACCCTTC 226

Search completed: Sun Oct 24 17:50:39 1999
Job time : 475 secs.

Query Match	12.48;	Score 29;	DB 12;	Length 114;
Best Local Similarity	4.98;	Pred. No. 8.13e-04;		
Matches	5;	Conservative	29;	Mismatches 69;
				Indels 0;
				Gaps 0;

RESULT	8
ID Q70468	standard; DNA; 114 BP.
Q70468	

	Location/Qualifiers
Key	55..60
misc-feature	/*tag a
	/note= "this sequence represents 'x'; 2 can be a
	sequence of 6, 9 or 12 nucleotides (see
	comments)"
05-APR-1995 (first entry)	
DE Generic DNA sequence to generate a random TSAR peptide library.	
KW TSAR: totally synthetic affinity reagent; synthetic; binding domain;	
KW effector domain; concatenated heterofunctional protein; linker;	
KW direct; rapid; detection; screening; treatment; generic; ss.	
OS Synthetic.	
FN Key	
FN misc-feature	
FN 55..60	
FN /*tag a	
FN /note= "this sequence represents 'x'; 2 can be a	
FN sequence of 6, 9 or 12 nucleotides (see	
FN comments)"	
PN N09A18318-A.	
PD 18-AUG-1994.	
PR 01-FEB-1994; U00977.	
PR 01-FEB-1993; US-013416.	
PR 30-DEC-1993; US-176500.	
PR 31-JAN-1994; US-189331.	
PA (UYNC-) UNIV NORTH CAROLINA.	
PI Fowles DM. Key BK;	
PI WPI: 94-279739/34.	
PT P-PSDB; R65154.	
PT Identifying proteins or peptide(s) which bind a ligand - by	
PT screening a recombinant vector library expressing fusion proteins	
PT comprising a binding domain and an effector domain	
PS Discourse; Page 35; 255pp. English.	
CC Q70468 is a generic DNA sequence used to generate random TSAR (Totally	
CC Synthetic Affinity Reagents) peptides. This generic formula can also be	
CC represented as follows: X(NNB)11(TGC)(NNB)62(NNB)7(TGC)(NNB)10Y. X	
CC and Y are flanking restriction sites (X is not the same as Y) that are	
CC not specified further. Other generic sequences are shown in Q70466-68.	
CC Other specific peptides generated by these generic sequences are shown in	

Query Match	12.0%;	Score 28;	DB 12;	Length 114;
Best Local similarity	2.7%;	Pred. No. 2.65e-03;		
Matches	3;	Conservative	31;	Mismatches 78; Indels 0; Gaps 0

RESULT	9
ID	Q70472 standard; DNA; 114 BP
NC	070473.

DT	10-APR-1995 (first entry)	
DE	Generic DNA sequence to generate a random TSAR peptide library.	
TSAR	totally synthetic affinity reagent; synthetic; binding domain;	
KW	effector domain; concatenated heterofunctional protein; linker;	
KV	direct; rapid; detection; screening; treatment; generic; ss.	
OS	synthetic.	
Key	Location/Qualifiers	
FT	misc-feature	55..60

FT /note= "encoded by 2"
 PM NC06183181-5.
 PD 18-AUG-1994.
 PF 01-FEB-1994; U00977.
 PR 01-FEB-1993; US-013416.
 PR 30-DEC-1993; US-176500.
 PR 31-JAN-1994; US-189331.
 PA (UYN-) UNIV NORTH CAROLINA.
 PI Fowlkes DM, Key BK;
 DR WPI: 94-279739/34.
 P-PSDB: RS6383.
 PT Identifying proteins or peptide(s) which bind a ligand - by
 PT screening a recombinant vector library expressing fusion proteins
 PS comprising a binding domain and an effector domain
 PS Disclosure: page 35; 25pp; English.
 CC Q70472 is a generic DNA sequence used to generate random TSAR (Totally
 CC Synthetic Affinity Reagents) peptides. This generic formula can also be
 CC represented as follows: X(NNB)1(CAC)(NNB)11(CAC)(NNB)(CAC)(NNB)22(CNNB)6
 CC -(CAC)(NNB)5(CAC)12(CNNB)4. X and Y are flanking restriction sites
 CC (X is not the same as Y) that are not specified further. The peptides
 CC generated by this and other generic sequences (Q70470-73) have invariant
 CC histidine residues incorporated into variant sequences. TSARs are
 CC concatenated heterofunctional proteins or peptides, comprising at least
 CC two functional regions - a binding domain with affinity for a ligand and
 CC a second effector peptide portion that is chemically or biologically
 CC active. They may further comprise a linker peptide between the 2 domains.
 CC The TSARs or compms, comprising a TSAR binding domain can be used in
 CC vivo to deliver a chemically or biologically active moiety, eg. metal
 CC ion, radioisotope, peptide, toxin or enzyme, to the specific target or
 CC on the cell. They can also replace the function of macromolecules, eg.

14.08; Score 28; DB 12; Length 114;

WO9418318-A

14.08; Score 28; DB 12; Length 114;

[illegible]

```

PR      Identifying proteins or peptide(s) which bind a ligand - by
PR      screening a recombinant vector library expressing fusion proteins
PT      Disclosures: Page 35; 25pp; English.
CC      Q70469 is a generic DNA sequence used to generate random TSAR peptide
CC      This generic formula can be represented as follows: X(TGGC)(NNB)10-
CC      T(GGC)(NNN)6(NNB)2(TTC)(NNB)14(TGCY). X and Y are flanking restriction
CC      sites (X is not the same as Y) that are not specified further. This
CC      sequence generates peptides that are cleavable in structure. Other
CC      genetic sequences are shown in Q70465-68. Other specific peptides
CC      generated by these functional proteins or peptides, comprising at least
CC      concatenated heterofunctional proteins or peptides, comprising at least
CC      two functional regions - a binding domain with affinity for a ligand and
CC      a second effector peptide portion that is chemically or biologically
CC      active. They may further comprise a linker peptide between the 2 domains.
CC      The oligonucleotides are also designed so that the expressed peptide
CC      contains 2 or 4 cysteine residues positioned in, or flanking, the
CC      unpredicted or variant residues. These residues confer some degree of
CC      conformational rigidity to the peptides. The TSARS or compns. comprising
CC      a TSAR binding domain can be used in vivo to deliver a chemically or
CC      biologically active moiety, eg. metal ion, radiotracer, peptide, toxin
CC      or enzyme, to the specific target or on the cell. They can also replace
CC      the function of macromolecules, eg. monoclonal or polyclonal antibodies
CC      and therefore circumvent the need for complex methods of hybridoma
CC      formation or in vivo antibody production. The TSARS are easily
CC      characterised and have designed activity allowing direct and rapid
CC      detection in a screening process.
SQ      Sequence 114 BP; 0 A; 4 C; 4 G; 4 T;
Query Match 11.1%; Score 26; DB 12; Length 114;
Best Local Similarity 1.98; Pred. No. 2,70e+02;
Matches 2; Conservative 30; Mismatches 74; Indels 0; Gaps 0;
Db        6 bnnbnbnbnbnbnbnbnbnbnbnbgcgnbnbnbnbnbnbnbnbnbnbnbnbnbnbn 65
Qy       134 GAGCTGAGGAGAAGTGTTTCTACAAATTCCTCATAGCGCACTTCGTGAATTGCT 193
          | . . . . . | . . . . . | . . . . . | . . . . . | . . . . .
Qy       194 TGAAAAGCTCTGCCCTCTCTCCATCTCCCTTAGGAGCACAGCGTC 239
          | . . . . . | . . . . . | . . . . . | . . . . . | . . . . .

RESULT   14
ID       Q51787 standard; DNA: 39 BP.
AC       Q51787;
DR       DTX_20-DXC-1993 (first entry)
DE       Mixed oligonucleotide #19 encodes ballast constituent.
KW       Fusion protein; ballast constituent; pro-insulin production;
KM       recombinant protein production; HMG CoA reductase;
KK       human 3-hydroxy-3-methylglutaryl-coenzyme A-reductase;
KN       mixed oligonucleotide; ss.
OS       Synthetic.
FH       Key Location/Qualifiers
FT       repeat_unit 4..6
FT                                     /*tag= a
FT /rpl_type=tandem
FT /note='can be repeated y times, where y is 4-11,
FT provided that y+z is 6-12'
FT 4..36
FT /*tag= b
FT /note="(DDP)11"
FT 35..38
FT /*tag= c
FT /rpl_type=tandem
FT /note='can be repeated z times, where z is 1-4,
FT provided that y+z is 6-12; N stands for
FT identical or different nucleotides,
FT excluding stop codons'
PN       US5227293-A.
PD       13-JUL-1993.
PF       29-AUG-1989; 399874.
PR       29-NUC-1989; US-399874.
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Query Match          12 0%:   Score 28; DB 3; Length 965;
Best Local Similarity 16 3%;
Matches      26; Conservative    65; Mismatches 69; Indels    0; Gaps    0;

Db      771 AANTGGCRKGDURHUVHVGCVRSTGTCTASDTTTSYWGWRGGMWDYGQTYNNGKR 830
        || : | : : : | : | : : | : | : : | : | : : | : | : : | : | : :
Cc      225 AAMGGAGATGAGCAGAGAGCGACACTTTTCACGCAATTCCAGAGACTCATGAG 166

Db      831 GRVTADPSSRSRYTADTAIVYYCVVSGSDGDDYWGTVYTSSHYUKMDITSSS 890
        || : | : : : | : | : : | : | : : | : | : : | : | : : | : | : :
Cc      165 AGGAATTTTGCGAAAAAGACTCTCTCACACTTCAGTGTGCAGAAGTATACCGGCCCTGTTC 106

Db      891 ASVGDRVYTCRSSSTTHGNNTYTWYTKGARYVSNRSRGVS 930
        || : : : : | : : : : | : : : : | : : : : | : : : : | : : : :
Cc      105 CATGTGACGTCCCAAGAGAGGCTCAAGAGAGGACAAGCGAGCGG 66

RESULT
ID US-08-238-163-5 STANDARD; DNA; UNC; 215 BP.
AC xxxxxx
DE
DN
Sequence 5, Application US/08238163
Cc Sequence 5, Application US/08238163
Cc Patent NO. 3569830
Cc GENERAL INFORMATION:
Cc APPLICANT: BENNETT, Alan
Cc APPLICANT: LABAVITCH, John M.
Cc APPLICANT: POWELL, Ann
Cc APPLICANT: STORZ, Henrik
Cc TITLE OF INVENTION: PLANT INHIBITORS OF FUNGAL
Cc TITLE OF INVENTION: POLYGALACTURONASES AND THEIR USE TO CONTROL FUNGAL DISEASE
Cc NUMBER OF SEQUENCES: 24
Cc CORRESPONDENCE ADDRESS:
Cc ADDRESSEE: Townsend and Townsend Kourlie and Crew
Cc STREET: Steuart Street Tower, One Market Plaza
Cc CITY: San Francisco
Cc STATE: California
Cc COUNTRY: US
Cc ZIP: 94105-1493
Cc COMPUTER READABLE FORM:
Cc MEDIUM TYPE: floppy disk
Cc COMPUTER: IBM PC compatible
Cc OPERATING SYSTEM: PC-DOS/MS-DOS
Cc SOFTWARE: PatentIn Release #1.0, Version #1.25
Cc CURRENT APPLICATION DATA:
Cc APPLICATION NUMBER: US/08/238,163
Cc FILING DATE: 03-MAY-1994
Cc CLASSIFICATION: 800
Cc ATTORNEY/AGENT INFORMATION:
Cc NAME: Bastian, Kevin L.
Cc REGISTRATION NUMBER: 34,774
Cc REFERENCE/DOCKET NUMBER: 2307E-540
Cc TELECOMMUNICATION INFORMATION:
Cc TELEPHONE: (415) 543-9600
Cc TELEFAX: (415) 543-5043
Cc INFORMATION FOR SEQ ID NO: 5:
Cc SEQUENCE CHARACTERISTICS:
Cc LENGTH: 215 base pairs
Cc TYPE: nucleic acid
Cc STRANDEDNESS: single
Cc TOPOLOGY: unknown
Cc MOLECULE TYPE: protein
Cc FEATURE:
Cc NAME/KEY: misc_feature
Cc LOCATION: 1..215
Cc OTHER INFORMATION: /standard_name="Deduced amino acid
Cc OTHER INFORMATION: sequence of PCIP from bean."
Cc SEQUENCE 215 BP; 15 A; 8 C; 25 G; 26 T; 141 OTHER.
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Db      18  CBNKAKXGNTSSMTWTCQCRNFNGCDPDTYVYNNDSGHAKKSSANTVCGANNVCAAK 77
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Qy      89  CCTTGACAGCTACATGACGAAAGGCCGGGATGATACCTTGCAAGCTGAACTG-AAAGAGT 147
      ::::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
Db      78  THYTHHWNSGADSKTYDYSYNASGTSNSGCTDGNRSGADSYGSSKNTAM 127
      ::::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
Qy      148  CTTTCTGACAAATTCCTCCATGAGTCCAGCTTCTCGGAAATGCTTGAA 197

RESULT 5
ID      US-08-676-967-2 STANDARD; DNA; UNC; 2277 BP.
AC      xxxxxx
DT
DE      Sequence 2, Application US/08676967
CC      Sequence 2, Application US/08676967
CC      Patent No. 5747317
CC      GENERAL INFORMATION:
CC      APPLICANT: COLLINS, KATHLEEN
CC      TITLE OF INVENTION: Human Telomerase
CC      NUMBER OF SEQUENCES: 10
CC      CORRESPONDENCE ADDRESS:
CC      ADDRESSER: Science & Technology Law Group
CC      STREET: 268 Bush Street, Suite 3200
CC      CITY: San Francisco
CC      STATE: CA
CC      COUNTRY: USA
CC      ZIP: 94104
CC      COMPUTER READABLE FORM:
CC      MEDIUM TYPE: Floppy disk
CC      COMPUTER: IBM PC compatible
CC      OPERATING SYSTEM: PC-DOS/MS-DOS
CC      SOFTWARE: Patentin Release #1.0, Version #1.30
CC      CURRENT APPLICATION DATA:
CC      APPLICATION NUMBER: US/08/676,967
CC      FILING DATE:
CC      CLASSIFICATION: 530
CC      ATTORNEY/AGENT INFORMATION:
CC      NAME: Osman Ph.D., Richard A
CC      REGISTRATION NUMBER: 36,627
CC      REFERENCE/DOCKET NUMBER: USB96-055
CC      TELECOMMUNICATION INFORMATION:
CC      TELEPHONE: (415)343-4341
CC      TELEFAX: (415)343-4342
CC      INFORMATION FOR SEQ ID NO: 2:
CC      SEQUENCE CHARACTERISTICS:
CC      LENGTH: 2277 base pairs
CC      TYPE: nucleic acid
CC      STRANDEDNESS: double
CC      TOPOLOGY: linear
CC      MOLECULE TYPE: cDNA
CC      SEQUENCE 2277 BP; 511 A; 212 C; 395 G; 216 T; 943 OTHER.

Query Match 9.8%; Score 23; DB 2; Length 2277;
Best Local Similarity 40.0%; Pred. No. 5,686-02;
Matches 18; Conservative 13; Mismatches 14; Indels 0; Gaps 0;

Db      1036  WSNAGARGARGARATYNGCAGARTATNTCAGCARTTGGCNGAR 1080
      ::||::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Cc      224  AGCGATGATGAGAGGAGCGACGATCTTTCACAGCAATTCACGAGAA 180

RESULT 6
ID      US-08-676-974-2 STANDARD; DNA; UNC; 2277 BP.
AC      xxxxxx
DT
DE      Sequence 2, Application US/08676974
CC      Sequence 2, Application US/08676974
CC      Patent No. 5770422
CC      GENERAL INFORMATION:
CC      APPLICANT: COLLINS, KATHLEEN
CC      TITLE OF INVENTION: Human Telomerase
CC      NUMBER OF SEQUENCES: 10

```

CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Science & Technology Law Group
CC STREET: 268 Bush Street, Suite 3200
CC CITY: San Francisco
CC STATE: CA
CC COUNTRY: USA
CC ZIP: 94104
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC OPERATING SYSTEM: IBM PC compatible
CC SOFTWARE: Patent Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/676,974
CC FILING DATE:
CC CLASSIFICATION: 530
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Osman P.D., Richard A
CC REGISTRATION NUMBER: 36,627
CC REFERENCE/DOCKET NUMBER: UCB96-055
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (415)343-4341
CC TELEFAX: (415)343-4342
CC INFORMATION FOR SEQ ID NO: 2:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 2277 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: double
CC TOPOLOGY: linear
CC MOLECULE TYPE: CDNA
CC SEQUENCE 2277 BP; 511 A; 212 C; 395 G; 216 T; 943 OTHER.

Query Match 9.8%; Score 23; DB 3; Length 2277;
Best Local Similarity 40.0%; Pred. No. 5,68e-02;
Matches 18; Conservative 13; Mismatches 14; Indels 0; Gaps 0;

Db 1036 WMSGARGARGARYTNGARGARYTNCARCATTYGCGNGAR 1080
Cp 224 AGCGAGATGAGGAGGAGGAGCACTTTCAGCAATTCAGCA 180

RESULT 7
ID PCT-US95-11934-99 STANDARD: DNA; UNC; 75 BP.
AC xxxxxx
DE Sequence 99, Application PC/TUS9511934
CC Sequence 99, Application PC/TUS9511934
CC GENERAL INFORMATION:
CC APPLICANT: Cytogen Corporation
CC TITLE OF INVENTION: Antigen Binding Peptides (Ablides) from
CC TITLE OF INVENTION: Peptide Libraries
CC NUMBER OF SEQUENCES: 103
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Pennie & Edmonds
CC STREET: 1155 Avenue of the Americas
CC CITY: New York
CC STATE: New York
CC COUNTRY: USA
CC ZIP: 10036
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC OPERATING SYSTEM: IBM PC compatible
CC SOFTWARE: Patent Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: PCT/US95/11934
CC FILING DATE: 20-SEP-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistrock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-196-228
CC TELECOMMUNICATION INFORMATION:

CC TELEPHONE: (212) 790-9090
CC TELEFAX: (212) 869-9741/8864
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 99:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 75 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: DNA (genomic)
CC SEQUENCE 75 BP; 1 A; 1 C; 7 G; 5 T; 61 OTHER.

Query Match 9.4%; Score 22; DB 4; Length 75;
Best Local Similarity 8.7%; Pred. No. 1.95e-01;
Matches 6; Conservative 19; Mismatches 44; Indels 0; Gaps 0;

Db 5 GANNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNN 64
Cy 54 GGCTCGGCGCTGCCCTTGTCTCTCTGACCTCTCTGCGACCTCATGACAGCG 113
Db 65 NBGCTGTG 73
Cy 114 CCGCGTAGG 122

RESULT 8
ID PCT-US95-11934-97 STANDARD: DNA; UNC; 82 BP.
AC xxxxxx

DE Sequence 97, Application PC/TUS9511934
CC Sequence 97, Application PC/TUS9511934
CC GENERAL INFORMATION:
CC APPLICANT: Cytogen Corporation
CC TITLE OF INVENTION: Antigen Binding Peptides (Ablides) from
CC TITLE OF INVENTION: Peptide Libraries
CC NUMBER OF SEQUENCES: 103
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Pennie & Edmonds
CC STREET: 1155 Avenue of the Americas
CC CITY: New York
CC STATE: New York
CC COUNTRY: USA
CC ZIP: 10036
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC OPERATING SYSTEM: IBM PC compatible
CC SOFTWARE: Patent Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: PCT/US95/11934
CC FILING DATE: 20-SEP-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistrock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-196-228
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (212) 790-9090
CC TELEFAX: (212) 869-9741/8864
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 97:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 82 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: DNA (genomic)
CC SEQUENCE 82 BP; 1 A; 2 C; 10 G; 8 T; 61 OTHER.

Query Match 9.4%; Score 22; DB 4; Length 82;
Best Local Similarity 8.7%; Pred. No. 1.95e-01;
Matches 6; Conservative 19; Mismatches 44; Indels 0; Gaps 0;

CC COUNTRY: U.S.A.
CC ZIP: 10036-2711
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patentin Release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/471,052A
CC FILING DATE: 06-JUNE-1995
CC CLASSIFICATION: 530
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistrock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-179
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: 212 790-9090
CC TELEFAX: 212 869-8864/9741
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 143:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 68 bases
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: unknown
CC MOLECULE TYPE: DNA
CC SEQUENCE 68 BP; 3 A; 3 C; 5 G; 3 T; 54 OTHER.

Query Match 9.0%; Score 21; DB 1; Length 68;
Best Local Similarity 13.8%; Pred. No. 6,54e-01;
Matches 9; Conservative 16; Mismatches 40; Indels 0; Gaps 0;

Db 4 CTGANNNA 63
Cp 178 CTGGACTCATAGAGGATTGTCAGAAAAGCTCCTGCTTCACTGCAAGTCA 119
Cc 64 CTTGG 68
Cc 118 CCCGG 114

RESULT 12
ID PCT-US95-11934-94 STANDARD; DNA; UNC; 74 BP.
AC xxxxxx
DE Sequence 94, Application PC/TUS9511934
CC Sequence 94, Application PC/TUS9511934
CC GENERAL INFORMATION:
CC APPLICANT: Cytogen Corporation
CC TITLE OF INVENTION: Antigen Binding Peptides (Abtides) From
CC NUMBER OF SEQUENCES: 103
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Pennie & Edmonds
CC STREET: 1155 Avenue of the Americas
CC CITY: New York
CC STATE: New York
CC COUNTRY: USA
CC ZIP: 10036
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patentin Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: PCT/US95/11934
CC FILING DATE: 20-SEP-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistrock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-196-228
CC TELECOMMUNICATION INFORMATION:

CC TELEPHONE: (212) 790-9090
CC TELEFAX: (212) 869-9741/8864
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 94:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 74 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: DNA (genomic)
CC SEQUENCE 74 BP; 3 A; 4 C; 3 G; 1 T; 63 OTHER.

Query Match 9.0%; Score 21; DB 4; Length 74;
Best Local Similarity 4.6%; Pred. No. 6,54e-01;
Matches 3; Conservative 19; Mismatches 43; Indels 0; Gaps 0;

Db 5 GANN 64
Oy 60 GGCGTCCCTTGTCTCTCTGACCTCCTTGACCTCAGTGAACAGGGCGGGT 119
Cc 65 NBNAC 69
Cc 120 ATGAC 124

RESULT 13
ID PCT-US95-11934-100 STANDARD; DNA; UNC; 74 BP.
AC xxxxxx
DE Sequence 100, Application PC/TUS9511934
CC Sequence 100, Application PC/TUS9511934
CC GENERAL INFORMATION:
CC APPLICANT: Cytogen Corporation
CC TITLE OF INVENTION: Antigen Binding Peptides (Abtides) From
CC NUMBER OF SEQUENCES: 103
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Pennie & Edmonds
CC STREET: 1155 Avenue of the Americas
CC CITY: New York
CC STATE: New York
CC COUNTRY: USA
CC ZIP: 10036
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patentin Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: PCT/US95/11934
CC FILING DATE: 20-SEP-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistrock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-196-228
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (212) 790-9090
CC TELEFAX: (212) 869-9741/8864
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 100:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 74 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: DNA (genomic)
CC SEQUENCE 74 BP; 6 A; 6 C; 1 G; 1 T; 60 OTHER.

Query Match 9.0%; Score 21; DB 4; Length 74;
Best Local Similarity 13.7%; Pred. No. 6,54e-01;
Matches 10; Conservative 17; Mismatches 46; Indels 0; Gaps 0;

US-09-092-296-7.rni

Page 7

RESULT	14	
ID	PCT-US95-11934-100	STANDARD; DNA; UNC; 74 BP.
AC	xxxxxx	

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DE Sequence 100, Application PC/TUS9511934
CC Sequence 100, Application PC/TUS9511934
CC GENERAL INFORMATION:
CC APPLICANT: Cytogen Corporation
CC TITLE OF INVENTION: Antigen Binding Peptides (Abtides) From
CC TITLE OF INVENTION: Peptide Libraries
CC NUMBER OF SEQUENCES: 103
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Pennie & Edmonds
CC STREET: 1155 Avenue of the Americas
CC CITY: New York
CC STATE: New York
CC COUNTRY: USA
CC ZIP: 10036
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Plentlin Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: PCT/US95/11934
CC FILING DATE: 20-SEP-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mirock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-196-228
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (212) 790-9090
CC TELEFAX: (212) 869-9741/8864
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 100:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 74 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: DNA (genomic)
CC SEQUENCE 74 BP: 6 A; 6 C; 1 G; 1 T; 60 OTHER.

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Query Match	9.0%;	Score 21;	DB 4;	Length 74;
Best Local Similarity	7.5%;	Pred. No. 6.54e-01;		
Matches	5;	Conservative	19;	Mismatches 43;
			Indels	0;
			Gaps	0;

[illegible]

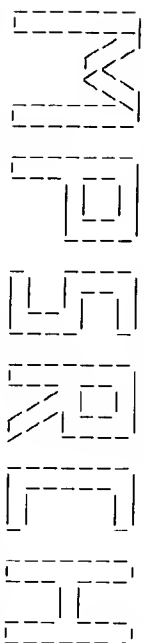
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RESULT 15
ID      PC/0595-11934-99 STANDARD; DNA; UNC; 75 BP
AC      XXXXXX
DT
DE      Sequence 99, Application PC/TUS9511934
CC      Sequence 99, Application PC/TUS9511934
GENERAL INFORMATION:

```

[illegible]

Search completed: Sun Oct 24 18:00:58 1999
Job time : 23 secs.



Release 3.1a John F. Collins, Biocomputing Research Unit.
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MSrch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 17:50:56 1999: MasPar time 454.93 Seconds

Tabular output not generated. 1205.245 Million cell updates/sec

Title: >US-09-092-296-7
Description: (51-284) from US09092296.seq
Perfect Score: 234
N.A. Sequence: 51 ATGGGCGTGGGCTGCCCCCT.....ACGATGTTGCTCAGACACA 284
Comp: TACCCGAGACCCGACGACGAGGGA.....TGGTACACAGACGCTGTGT

Scoring table: TABLE default

Gap 6

Mismatch STD : Dbase 0: Query 0

Searched: 2883791 seqs, 1171580779 bases x 2

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database:

embl-est58
1:em-est10 2:em-est11 3:em-est17 4:em-est18 5:em-est2
6:em-est9 7:em-gss1
Database: genbank-est111

13:gb-est14 9:gb-est10 10:gb-est11 11:gb-est12 12:gb-est13
17:gb-est18 18:gb-est15 15:gb-est16 16:gb-est17
21:gb-est21 22:gb-est19 19:gb-est20 20:gb-est24
25:gb-est25 26:gb-est26 27:gb-est27 28:gb-est28
29:gb-est29 30:gb-est31 31:gb-est32 32:gb-est33 33:gb-est34
34:gb-est35 35:gb-est36 36:gb-est37 37:gb-est38 38:gb-est39
39:gb-gss3 40:gb-gss4 41:gb-gss5 42:gb-gss6

Statistics: Mean 9.804; Variance 2.109; scale 4.649

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description	Pred. No.
1	73	31.2	328	23	AI136523	UT-R-C2P-ng-e-02-0-01.
2	57	24.4	252	17	AA154439	7.25e-84
3	41	17.5	252	17	AA154439	2.49e-56
4	33	14.1	247	17	AA154458	2.28e-30
5	33	14.1	247	17	AA154458	2.20e-18
6	32	13.7	2275	20	AF034173	5.89e-17
7	27	11.5	1287	20	AF038250	4.09e-10
8	27	11.5	2275	20	AF034173	8.11e-09
9	26	11.1	196	21	AF034173	4.09e-10
10	25	10.7	176	19	T94049	1.51e-07

C	11	10.7	398	8	T52782	va79f01.r1 Stratagene	1.51e-07
C	12	10.7	582	37	B04965	CSRL-51A6-u cSRL flow	1.51e-07
C	13	9.8	364	34	B33870	mc56b03.r1 Soares mous	4.22e-05
C	14	9.8	397	30	R52894	YH01009.r1 Soares Infa	4.22e-05
C	15	9.8	400	28	AI554565	tn25106.x1 NCI_CGAP-Br	4.22e-05
C	16	9.8	420	31	H56592	yt88b09.r1 Soares_pine	4.22e-05
C	17	9.8	426	8	T15255	cr5855_lamdaZAPst Ric	4.22e-05
C	18	9.8	459	13	AA475002	vh08h04.r1 Soares mous	4.22e-05
C	19	9.8	461	22	AI086444	qf25e01.x1 NCI_CGAP-Br	4.22e-05
C	20	9.8	547	25	AI337590	qf31a08.x1 NCI_CGAP-Br	4.22e-05
C	21	9.8	1287	20	AF038250	AF038250 Human mRNA (T	4.22e-05
C	22	9.4	201	28	AI353335	UT-R-C3-sw-e-12-0-UT.s	6.28e-04
C	23	9.4	259	38	B81136	CIT-HSP-2015F16.TFC CI	6.28e-04
C	24	9.4	308	8	D40392	RICS2342A Rice shoot O	6.28e-04
C	25	9.4	334	33	W84051	mt6e11.r1 Soares mous	6.28e-04
C	26	9.4	404	17	AA730403	n442c07.s1 NCI_CGAP-Ev	6.28e-04
C	27	9.4	408	17	T46345	9608_Lambda-Phl2 Arabi	6.28e-04
C	28	9.4	424	33	W13114	ma86e10.r1 Soares mous	6.28e-04
C	29	9.4	433	34	W79098	zd75h10.r1 Soares feta	6.28e-04
C	30	9.4	441	14	C28493	C28493 Rice callus CDN	6.28e-04
C	31	9.4	459	27	AI442028	ss66f12.y1 Gam-cl004 G1	6.28e-04
C	32	9.4	481	42	AQ445188	GSSTC01459 Trypanosoma	6.28e-04
C	33	9.4	513	40	AQ235722	HS-2015_B2_C08.T7 CIT	6.28e-04
C	34	9.4	602	23	AI165652	xy1em.est.742 Poplar x	6.28e-04
C	35	9.4	634	32	AI063013	GH02423.5prime GH Dros	6.28e-04
C	36	9.4	1023	37	B12587	F22011-Sp6.1 IGF Arabi	6.28e-04
C	37	9.0	317	19	F10052	HSC39H12 normalized 1	8.54e-03
C	38	9.0	354	26	AI398058	NCMA1117 Subtracted	8.54e-03
C	39	9.0	422	33	M07189	za94p07.r1 Soares_feta	8.54e-03
C	40	9.0	516	31	H41536	y771c01.s1 Soares adu1	8.54e-03
C	41	9.0	524	16	AA629071	tf88f08.s1 Soares-test	8.54e-03
C	42	9.0	550	20	AA898701	NCM667Y Mycelial Neur	8.54e-03
C	43	9.0	567	30	R61539	yh16f01.s1 Soares Infa	8.54e-03
C	44	9.0	587	13	AA428548	zw47d06.r1 Soares tota	8.54e-03
C	45	9.0	630	39	AQ201160	RPC111-46k18.TJ RPC111	8.54e-03

ALIGNMENTS

RESULT LOCUS	1	AI136523	328 bp	mRNA	EST	11-FEB-1999
DEFINITION		UT-R-C2P-ng-e-02-0-01.s1	UT-R-C2P	Rattus norvegicus	CDNA clone	
ACCESSION		AI136523				
NID		93637300				
VERSION		AI136523.1	GI:3637300			
KEYWORDS						
SOURCE						
ORGANISM						
REFERENCE						
AUTHORS						
TITLE						
JOURNAL						
MEDLINE						
COMMENT						

On Jan 14, 1998 this sequence version replaced gi:1877567.

Genome Res. 6 (9), 791-806 (1996)

97044477

Contact: Soares, MB

Program for Rat Gene Discovery and Mapping

University of Iowa

451 Eckstein Medical Research Building Iowa City, IA 52242, USA

Tel: 319 335 8250

Fax: 319 335 9555

Email: msoares@iue.uiowa.edu

The sequence tag present in the CDNA between the NotI site and the

oligo of track served to identify it as a clone from the normalized

adult lung library. CDNA Library Preparation: M. Fatima Bonaldo,

Ph.D. Clone distribution: clones will be available through Research

Genetics

Seq primer: M13 Forward.

Location/Qualifiers

```

1. 328
/organism="Rattus norvegicus"
/strain="Sprague-Dawley"
/vector="pTR70-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; The UI-R-C2p
library is a subtracted library derived from the UI-R-C1
library, which is a subtracted library derived from the
UI-R-C0 library. The UI-R-C0 library consisted of a
mixture of individually tagged normalized libraries
constructed from rat placenta, adult lung, brain, liver,
kidney, heart, spleen, ovary, muscle, 8, 12 and 18-day
embryo. The tag is a string of
3-5 nucleotides present
between the Not I site and the oligo-OT track which allows
identification of the library of origin of a clone within
the mixture. The subtracted library (UI-R-C2p) was
constructed as follows: PCR amplified cDNA inserts from
UI-R-C1 clones from which 3' ESTs had been derived was
used as a driver in a hybridization with the UI-R-C1
library in the form of single-stranded circles. The
remaining single-stranded circles (subtracted library) was
purified by hydroxapatite column chromatography
converted to double-stranded circles and electroporated
into DH10B bacteria (Life Technologies) to generate the
UI-R-C2p library. This procedure has been previously
described (Donaldd, Lennon and Soares, Genome Research 6:
791-806, 1996)".
/db_xref="taxon:10116"
/clone="UI-R-C2p-ng-e-02-0-UI"
/clone_1fb="UI-R-C2p"
/dev_string="adult"
/lab_host="db10b (Life Technologies)"
a 77 c 98 g 91 t

```

```

Department of Cytogenetics
National Inst. of Agril. Sci. and Tech, RDA
Suwon, Kyunggi-do, Korea
Tel : 82 331 290 0301
Fax: 82 331 290 0307
Email: myeungsun20.as@nrc.ac.kr
Submitted by Baek Hye Nahm, Dept of Biological Science, Myongji
University, Yongin, Korea. 449-728 bnhna@bioserver.myongji.ac.kr
Seq primer: M13 Reverse Primer.

FEATURES
        source
            location/Dualifiers
                1..252
                    /organism="Oryza sativa"
                    /cultivar="Milyang23"
                    /note="Vector: pBluescript SK(+); Site_1: EcoRI; Site_2:
XhoI; Directional cDNA library inserted into Lambda ZAP101
vector at 5' end with EcoRI and 3' end with Xho I site."
                    /db_xref="taxon:4530"
                    /map=6"
                    /clone="97SN187"
                    /clone_lib="Rice Immature Seed Lambda ZAPII cDNA Library"
                    /tissue_type="Immature Seed"
                    /dev_stage="5 days after pollination"
                    /adb_host="E. coli SOLR"
                    21 c          12 g          35 t          179 others

BASE COUNT
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Best Local Similarity 10.9%; Pred. No. 2.49e+56;
Matches 23; Conservative 108; Mismatches 78; Indels 2; Gaps 2;

DB 22 SYBAGNBNWVVAASHGNGVMYHNCETRGTHCDCKNVNMSTMGTCTNNBNYSQDMHYNB 81
::: :::: |:::: ::::: ::::: ::::: ::::: ::::: ::::: ::::: ::::: :::::
49 CTTCGTCCTCCTTGAACCTCCTTGCGATCATGCATGCAGACAGGCCGGTGTACTTTG 128

```

Query Match	31.2%	Score 73	DB 23	Length 328
Best Local Similarity	73.0%	Pred. No. 7,25e-64		
Matches 116	Conservative 0	Mismatches 43	Indels 0	Gaps 0
Db 142	TCGACGTAAGGTGTGTGTGGTGGCCCTTTATGATGAAGGAGTGACATTGTGCCCGGTGAGA	201		
Cp 279	TCGACACAACTGCTGTGAATCTTCAATGATGGAGGAGTACCGCTGCCCTCAGAGCA	220		
Db 202	GGTGAGAGAGAGACAGACATCTTTGGAGCAATGCCAAGAACCGGAGCTTGGAGAGTCT	261		
Cp 219	GATGAGAGAGAGAGCAGACCTTTTCAAGCAATTCAGGAAGCTGGACATAGAGAGCAAT	160		
Db 262	TGGCTGAGATGCCGTGGCTGACATTTTACCTTCAAAATGCA	300		
Cp 159	TTGTCAGAAAAGACTCTTCAGCTTCAGTTCCAAAGTCA	121		
RESULT 2				
LOCUS	AA754459	252 bp	mRNA	EST
DEFINITION	97SN1787 Rice Immature Seed Lambda ZAP1	CDNA	Library	Oriza sativa
ACCESSION	AA754459			
NID	92801165			
VERSION	AA754459.1	GI:2801165		
KEYWORDS	EST.			
SOURCE	Oriza sativa.			
ORGANISM	Oriza sativa.			
REFERENCE	Eukaryotes: Vitidiplantae: Streptophyta: Embryophyta: Tracheophyta: eumphylophytes: Spermatophyta: Magnoliophyta: Liliopsida: Poales; Poaceae: Oryza.			
AUTHORS	1 (bases 1 to 252)			
TITLE	Nahn,B.J., Kim,J.K., Cheong,J.J., Kim,S.I., Hahn,T.R., Moon,E.P., Kim,W.T., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y., Lee,M.C. and Eun,M.Y.			
JOURNAL	Large-scale Sequencing Analysis of ESTs from Rice Immature Seed Unpublished (1998)			
COMMENT	On Jan 14, 1998 this sequence version replaced gi:1197457.			
Contact:	Eun M.Y.			

RESULT	3	252 bp	mRNA	EST	20-JAN-1998
LOCUS	AA754459				
DEFINITION	97SN1187 Rice Immature Seed Lambda ZAPRI cDNA Library Oryza sativa				
ACCESSION	AA754459				
NID	92801165				
VERSION	AA754459.1	GI:2801165			
KEYWORDS					
SOURCE	EST.				
ORGANISM	Oryza sativa.				
	Oryza sativa				
	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;				
	euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;				
	Poaceae; Oryza.				
REFERENCE	1 (bases 1 to 252)				
AUTHORS	Nahm,B.H., Kim,J.K., Cheong,J.J., Kim,S.I., Hahn,T.R., Moon,E.P.,				
	Kim,W.T., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y.,				
	Lee,M.C. and Eun,M.Y.				
TITLE	Large-Scale Sequencing Analysis of ESTs from Rice Immature Seed				
JOURNAL	Unpublished (1998)				
COMMENT	On Jan 14, 1998 this sequence version replaced gi:1179457.				

Contact: Eun M.Y.
 Department of Cytogenetics
 National Inst. of Agri. Sci. and Tech, RDA
 Suwon, Kyunggi-do, Korea
 Tel: 82 331 250 0301


```
BASE COUNT      7 a      16 c      21 g      34 t      169 others
ORIGIN
Query Match      14.1%; Score 33; DB 17; Length 247;
Best Local Similarity 14.7%; Pred. No. 2.20e-18;
Matches 24; Conservative 71; Mismatches 66; Indels 2; Gaps 1;

Db 34 VRRVGTNNKNGHRTTMMDCSDNACRYTBYMYANSKYGYGTBYYSKNVDNTTGT 93
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 223 GGGAGATGGAGGAGGAGGAGCTTTTCAAGCAATTCAGAACTGACTATAGGAAT 164
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 94 GYCKTTVAVHSGMNRCSNVYVMBTAYCDYBHDNRNHYDDRCINDRGCONTASD 153
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 163 GAATTTGTCAAGAAAGACTCTTCAGCTTCAGTTCACCAAGTCAACCGGCTCTTCCA 104
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 154 NGTSATKRYTGYDKTSDCGGCKRYTYGSSBYBRGVNVAV 196
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 103 TGTGA--GCTGCCAAGGAGGAGGCAAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 63
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 6 AF034173 2275 bp mRNA EST 30-MAR-1998
LOCUS AF034173 Human mRNA (Tripodis and Ragousis) Homo sapiens cDNA
DEFINITION Clone ntcon2 contig, mRNA sequence.
ACCESSION AF034173
NID 92707735
VERSION AF034173.1 GI:2707735
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 2275)
AUTHORS Tripodis,N. and Ragousis,J.
TITLE Generation of a transcription map in the region immediately
centromeric to human MHC across the 6p21.2-6p21.3 chromosomal
boundary
JOURNAL Unpublished (1997)
COMMENT On Jan 19, 1998 this sequence version replaced gi:2045115.

Contact: Tripodis, Nikos
Division of Medical and Molecular Genetics
Guys Hospital
7th floor, Guy's Tower, London SE1 9RT, UK
Email: nikos@nki.ni.

FEATURES
Source
1..2275
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="6p21.3"
/clone="ntcon2 contig"
/clone_11b="Human mRNA (Tripodis and Ragousis)"
BASE COUNT 438 a 619 c 470 g 599 t 149 others
ORIGIN
Query Match      13.7%; Score 32; DB 20; Length 2275;
Best Local Similarity 11.2%; Pred. No. 5.89e-17;
Matches 13; Conservative 66; Mismatches 35; Indels 2; Gaps 2;

Db 1480 RYKRMKRRKRRKRRMTGMYRFRMMAMCAMACMYWYWKRMKMKCKRYKRYKYS 1539
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 218 ATGAGAGAGGAGGAGGAGCTTTTCAAGCAATTCAGAACT--GACTCATAGAGGAAT 160
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1540 TTYASMSRMTWTTTYYTCWCT--SKASASAMRMKGYGSSRSRSTWGTGNSM 1594
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 159 TTGTCAAGAAAGACTCTTCAGCTTCAGTTCAGTTCACCAAGTCAACCGGCTCTTCCA 104
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 7 AF038250 1287 bp mRNA EST 30-MAR-1998
LOCUS AF038250
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AF038250 Human mRNA (Tripodis and Ragousis) Homo sapiens cDNA
DEFINITION Clone ntcon3, mRNA sequence.
ACCESSION AF038250
NID 92815880
VERSION AF038250.1 GI:2815880
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Tripodis,N. and Ragousis,J.
TITLE Generation of a transcription map in the region immediately
centromeric to human MHC across the 6p21.2-6p21.3 chromosomal
boundary
JOURNAL Unpublished (1997)
COMMENT On Jan 19, 1998 this sequence version replaced gi:2045085.

Contact: Tripodis, Nikos
Division of Medical and Molecular Genetics
Guys Hospital
7th floor, Guy's Tower, London SE1 9RT, UK
Email: nikos@nki.ni.

FEATURES
Source
1..1287
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="6p21.3"
/clone="ntcon3"
/clone_11b="Human mRNA (Tripodis and Ragousis)"
BASE COUNT 349 a 219 c 293 g 361 t 65 others
ORIGIN
Query Match      11.5%; Score 27; DB 20; Length 1287;
Best Local Similarity 45.6%; Pred. No. 4.09e-10;
Matches 26; Conservative 15; Mismatches 16; Indels 0; Gaps 0;

Db 432 GRYGVBGASTCTMYNCCDCKTGSAGVTVNHHDSNAGAGAAAGCTTCTTCGCA 488
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 246 GGGAGGTGACGCTGTCCTCGAAGGAGATGAGGAGGAGGAGGAGGAGGAGGAGGAG 190
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 8 AF034173 2275 bp mRNA EST 30-MAR-1998
LOCUS AF034173 Human mRNA (Tripodis and Ragousis) Homo sapiens cDNA
DEFINITION Clone ntcon2 contig, mRNA sequence.
ACCESSION AF034173
NID 92707735
VERSION AF034173.1 GI:2707735
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 2275)
AUTHORS Tripodis,N. and Ragousis,J.
TITLE Generation of a transcription map in the region immediately
centromeric to human MHC across the 6p21.2-6p21.3 chromosomal
boundary
JOURNAL Unpublished (1997)
COMMENT On Jan 19, 1998 this sequence version replaced gi:2045115.

Contact: Tripodis, Nikos
Division of Medical and Molecular Genetics
Guys Hospital
7th floor, Guy's Tower, London SE1 9RT, UK
Email: nikos@nki.ni.

FEATURES
Source
1..2275
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="6p21.3"
/clone="ntcon2 contig"
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polylinker. Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5'] TGTTACCAATCGAGTGGCGGCGCGGCAATTTTCTTTTCTTTT T 3']. on equal amounts of mRNA from 2 13.5dpc and 2 14.5dpc embryos [total RNA provided by Minoru Ko, Wayne State Univ., from 2]; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo.

/db_xref="taxon:10090"
/clone_image="352493"
/clone_lib="Soares mouse embryo NDME13.5 14.5"
/sex="unknown"
/tissue_type="embryo"
/dev_stage="13.5-14.5dpc total fetus"
/lab_host="DH10B"

BASE COUNT

100 a 73 c 87 g 104 t

ORIGIN

Query Match 9.8%; Score 23; DB 34; Length 364;
Best Local Similarity 79.5%; Pred. No. 4.22e-05;
Matches 31; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

Db 282 AACCTGAAGGAGGAGGCTTCCACATATTTCTCT 320
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OY 130 AACTGAGCTGAGAGCTTTTTCGCAATTCCTCT 168

RESULT 14

LOCUS R52894 397 bp mRNA EST 18-MAY-1995
DEFINITION y01d09.r1 Soares Infant Brain INIB Homo sapiens cDNA clone
IMAGE:41873 5', mRNA sequence.

ACCESSION R52894
NID 9814796
VERSION R52894.1 GI:814796
KEYWORDS EST.
SOURCE human

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 397)

REFERENCE

AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Mente, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaaskis, E., Waterston, R., Williamson, A., Wohlmann, P., and Wilson, R.
The WashU-Merck EST Project
Unpublished (1995)

TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT On Sep 21, 1992 this sequence version replaced gi:275991.

Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: estewatson.wustl.edu

High quality sequence stops: 299
Source: IMAGE Consortium, LNL
This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information.
Seq primer: M13RP1
High quality sequence stop: 299.
Location/Qualifiers

FEATURES

Source

/organism="Homo sapiens"
/note="Organ: whole brain; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5'] TGTTACCAATCGAGTGGCGGCGGCAATTTTCTTTTCTTTT T 3']; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and Eco RI sites of the modified pT73 vector. Library is normalized, and was constructed by Bento Soares and M. Fatima Bonaldo."
/db_xref="taxon:9606"
/map="19p12-p13.1: p12"
/clone_image="2168675"
/clone_lib="NCL_CGAP_Brn25"

into the Not I and Hind III sites of the pT73D-Pac vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo.

BASE COUNT 115 a 74 c 100 g 104 t 4 others

ORIGIN

Query Match 9.8%; Score 23; DB 30; Length 397;
Best Local Similarity 78.0%; Pred. NO. 4.22e-05;
Matches 32; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

Db 7 GAGGAGAGTCAAGACATTCAGATTTCCAGAGAGTGC 47
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Cp 215 GAGGAGAGGAGGAGGCTTTTCAGCAATTCAGAGAGCTGC 175

RESULT 15

LOCUS A1554565 400 bp mRNA EST 23-MAR-1999
DEFINITION t25t06.x1 NCL_CGAP_Brn25 Homo sapiens cDNA clone IMAGE:2168675 3', similar to TR:Q16664 Q16664 PROTEIN A-1. [1] ;, mRNA sequence.

ACCESSION A1554565
NID 94486928
VERSION A1554565.1 GI:4486928
KEYWORDS EST.
SOURCE human

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 400)

REFERENCE

AUTHORS NCI/NINDS-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute / National Institute of Neurological Disorders and Stroke, Brain Tumor Genome Anatomy Project (CGAP/BRGAP), Tumor Gene Index
Unpublished (1998)
On May 7, 1998 this sequence version replaced gi:3121680.

JOURNAL COMMENT

Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: www-bio.lnl.gov/dbp/image/image.html

Seq primer: -400P from Gibco.
Location/Qualifiers

FEATURES

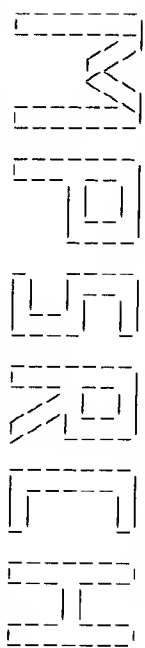
Source

/organism="Homo sapiens"
/note="Organ: Brain; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5'] TGTTACCAATCGAGTGGCGGCGGCAATTTTCTTTTCTTTT T 3']; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and Eco RI sites of the modified pT73 vector. Library is normalized, and was constructed by Bento Soares and M. Fatima Bonaldo."
/db_xref="taxon:9606"
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/tissue_type="anaplastic oligodendroglioma"
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 Best Local Similarity 74.5%: Pred. No. 4.22e-05;
 Matches 35; Conservative 0; Mismatches 12; Indels 0; Gaps 0;
 DB 268 AAGCCTTGCACTTTCTTCACCCCGTCAGGAGAGAGCTCTCCCT 314
 QY 198 AAGCTTGCCCTCCTCCATCTCCCTTCAGGAGAGAGCTACCCCT 244

Search completed: Sun Oct 24 17:58:39 1999
 Job time : 463 secs.



(TM)

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Maspar_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 17:19:23 1999; Maspar time 370.71 Seconds

Tabular output not generated. 1345,905 Million cell updates/sec

Title: >US-09-092-296-3
Description: (1-180) from US0902296.seq
Perfect Score: 180
N.A. Sequence: 1 CAGGAGGCGAGTGGCCACTA.....CTTGAAGACTCTCTCCCTCT 180
Comp: GTCCTCCGCTCACCGCTCAT.....GACCTTTGAGACGAGAGA

Scoring table: TABLE default
Gap 6

Mismatch STD: Dbase 0; Query 0

Searched: 646147 seqs, 1385953633 bases x 2

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database: emb158
1:em_ba1 2:em_ba2 3:em_fun 4:em_htg 5:em_hum1 6:em_hum2
7:em_in 8:em_com 9:em_or 10:em_cov 11:em_pat 12:em_ph
13:em_pl 14:em_ro 15:em_sts 16:em_vi
Database: genbank111
17:gb_ba1 18:gb_ba2 19:gb_hgt1 20:gb_hgt2 21:gb_in1
22:gb_in2 23:gb_com 24:gb_cov 25:gb_pat 26:gb_ph 27:gb_pl1
28:gb_pl2 29:gb_pl3 30:gb_pl4 31:gb_pl5 32:gb_ro
33:gb_st 34:gb_sts 35:gb_sy 36:gb_un 37:gb_vi

Statistics: Mean 9.304; Variance 4.612; scale 2.018

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description	Pred. No.
1	90	50.0	47323	31	AC005937	Homo sapiens clone UMG 8.42e-49
2	32	17.8	216021	31	HUAC004787	Homo sapiens Chromosome 7.20e-06
3	29	16.1	965	25	AR024229	Sequence 22 from patent 5.53e-04
4	28	15.6	7218	25	166494	Sequence 14 from patent 2.26e-03
5	27	15.0	215	25	128278	Sequence 5 from patent 9.05e-03
6	27	15.0	215	25	128278	Sequence 5 from patent 9.05e-03
7	26	14.4	965	25	AR024229	Sequence 22 from patent 3.54e-02
8	26	14.4	74371	31	AC005369	Homo sapiens Chromosome 3.54e-02
9	26	14.4	216021	31	HUAC004787	Homo sapiens Chromosome 3.54e-02
10	25	13.9	565	25	E04076	gDNA encoding envelope 1.35e-01
11	25	13.9	60966	31	AC003030	Homo sapiens chromosome 1.35e-01
12	24	13.3	60	25	A62889	Sequence 1 from patent 5.03e-01
13	24	13.3	1056	23	MV087256	Mustela vison GT dinuc 5.03e-01

14	23	12.8	1056	23	MV087256	Mustela vison GT dinuc 1.82e+00
15	23	12.8	390	29	HU06FUCAS	H. sapiens fucosidase p 1.82e+00
16	23	12.8	175902	29	AC006992	Homo sapiens clone NH0 1.82e+00
17	22	12.2	30	25	A62994	Sequence 6 from patent 6.36e+00
18	22	12.2	108	21	D87227	Trypanosoma cruzi mRNA 6.36e+00
19	22	12.2	1738	29	HU06FUCAS	Human (clone SY2/10) g 6.36e+00
20	22	12.2	1945	24	CHCONN45	Chicken connexin-45 mr 6.36e+00
21	22	12.2	180	21	DROSTY	D. melanogaster synapto 6.36e+00
22	22	12.2	2273	32	MBAKEIN	Mus musculus Bak gene, 6.36e+00
23	22	12.2	3088	11	E10775	DNA encoding part of A 6.36e+00
24	22	12.2	3088	11	E10775	DNA encoding part of A 6.36e+00
25	22	12.2	6640	29	D63997	Homo sapiens mRNA for 6.36e+00
26	22	12.2	67919	29	AC006542	WORKING DRAFT SEQUENCE 6.36e+00
27	22	12.2	82098	31	AC006252	Homo sapiens 3p21.1 co 6.36e+00
28	22	12.2	124700	22	AC005558	Drosophila melanogaste 6.36e+00
29	22	12.2	133457	30	AC003999	Human PAC clone DJ1139 6.36e+00
30	22	12.2	135039	31	AC006060	Homo sapiens 3p22-8 PA 6.36e+00
31	22	12.2	151187	19	HS4608	Human DNA sequence *** 6.36e+00
32	22	12.2	175339	31	AC005772	Homo sapiens chromosome 6.36e+00
33	22	12.2	203418	19	AC004947	Homo sapiens clone D11 6.36e+00
34	22	12.2	241407	20	AC003059	Mouse chromosome 10 BA 6.36e+00
35	21	11.7	65	25	I41365	Sequence 145 from pate 2.16e+01
36	21	11.7	631	30	HSATP2A152	Human Ca2+ ATPase of f 2.16e+01
37	21	11.7	1000	30	HSATP2A152	H. sapiens tal-1 DNA. 2.16e+01
38	21	11.7	1755	29	HS277017	Homo sapiens mRNA for 2.16e+01
39	21	11.7	3425	29	AB006623	Homo sapiens mRNA for 2.16e+01
40	21	11.7	46213	28	SPBC18H10	S. pombe chromosome II 2.16e+01
41	21	11.7	56804	30	HS77N19	Human DNA sequence fro 2.16e+01
42	21	11.7	74371	31	AC005369	Homo sapiens chromosome 2.16e+01
43	21	11.7	118995	31	AC005368	Homo sapiens chromosome 2.16e+01
44	21	11.7	127027	30	HS461P17	Human DNA sequence fro 2.16e+01
45	21	11.7	187966	31	AC006487	Homo sapiens chromosome 2.16e+01

ALIGNMENTS

RESULT 1
LOCUS AC005937 47323 bp DNA PRI 05-NOV-1998
DEFINITION Homo sapiens clone UMG:370M23.002 from 6p21, complete sequence.
ACCESSION AC005937
NID G3845393
VERSION AC005937.1 GI:3845393
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 47323)
AUTHORS Janer M., Guillaudoux, T., Vu, Q., Kucyavin, T., Harter, H. and Geraghty, D. E.
TITLE Large scale sequence analysis of the human MHC class I region
JOURNAL Unpublished (1998)
REMARK Fred Hutchinson Cancer Research Center
The Clinical Research Division
1100 Fairview Ave. N., P.O. Box 19024
Seattle, WA 98109-1024
2 (bases 1 to 47323)
Geraghty, D. E. and Olson, M. V.
Direct Submission
Submitted (05-NOV-1998) Human Genome Center, University of
Washington, Box 352145, Seattle, WA 98195, USA
University of Washington Human Genome Center
Box 352145 Seattle, WA 98195
Contact: Daniel E. Geraghty (geraghty@fhcr.org)
Overlaping Sequences:
5'-UMGC:370M23.013 (Genbank Accession: AC005530)
3'-UMGC:y67C112 (Genbank Accession: AC004211)

Sequence Quality Assessment:
This entry has been annotated with sequence quality estimates computed by the Phrap assembly program. All manually edited bases have been reduced to quality zero. Quality levels above 40 are expected to have less than

1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the
GenBank flat file format but are available as part
of this entry's ASN.1 file.

Double stranded (DS) coverage: 75.5%
DS or two chemistry coverage: 98.9%
Single stranded regions: 3

Sequence Validation:
This sequence has been validated by Multiple Complete Digest
Mapping. Comparison of the experimentally derived map digest
fragments with sequence-predicted fragments is given below.
Small fragments below a variable cutoff (approximately 400-600bp)
are not mapped and hence do not appear in the table. There are no
significant remaining discrepancies between the experimental and
predicted values. Uniquely ordered fragment groups are separated
by dashed lines.

Map	Seq	Map	Seq	Map	Seq
BglII		HindIII		NsiI	
1069.11	1050.00	889.55	866.00	30541.40	30653.00
20320.67	20855.00	1050.18	1015.00	3279.08	3231.00
2171.50	2147.00	7268.78	7196.00		
2560.20	2531.00	10085.80	9992.00		
4335.42	4269.00	11212.78	11131.00		
2698.62	2628.00				
1927.50	1887.00				
3130.46	3090.00				
2166.69	2129.00				
2044.67	2005.00				

FEATURES

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/db_xref="taxon:9606"
/chromosome="6"
/map="6p21"
/sub_clone="UMGC:370M23.002"
/clone_idb="Research Genetics BAC Library"
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complement(4999..5277)
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6285..6572
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25510..25802
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31295..31594
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33515..33767
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repeat_region 37372..37648
/rpt_family="Alu"
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/rpt_family="MER3"
repeat_region 39583..40010
/rpt_family="Alu"
repeat_region 40046..40156
/rpt_family="Alu"
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43325
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46851
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47032
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47240..47256
variation /note="clonal variation with 3' overlapping clone -
insertion of 17bp repeat"

BASE COUNT 11556 a 11489 c 12284 g 11994 t
ORIGIN

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Best Local Similarity 98.9%; Pred. No. 8,42e-49;
Matches 91; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Db 35465 CAGGGCGGATGACTCTTGCAAGTGAAGAGGAGCTTTTGTGCAAAATTCCTCT 35524
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Gy 78 CAGGGCGGATGACTCTTGCAAGTGAAGAGGAGCTTTTGTGCAAAATTCCTCT 137
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Db 35525 ATGAGTCCAGCTTCCTGGAATTCCTGGAAG 35556
|||||
Gy 138 ATGAGTCCAGCTTCCTGGAATTCCTGGAAG 169
|||||

RESULT 2
LOCUS HUAC004787 216021 bp DNA PRI 24-JUL-1998
DEFINITION Homo sapiens chromosome 16 BAC clone C1987SK-A-952F10, complete
sequence.
ACCESSION AC004787
NID 93337381
VERSION AC004787.1 GI:3337381
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Primates; Catarrhini; Hominoidea; Homo.
REFERENCE 1 (bases 1 to 216021)
AUTHORS Adams M.D., Loftus B.J., Zhou L., Crosby M., Fuhmann J.,
Mason T.M., Brandon R., Kim U.J., Keilavage A.R. and Venter J.C.
TITLE Homo sapiens chromosome 16 BAC clone C1987SK-A-952F10
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 216021)
AUTHORS Adams M.D. and Loftus B.J.
TITLE Direct Submission

REFERENCE 1 (bases 1 to 215)
AUTHORS Bennett,A., Labavitch,J.M., Powell,A. and Stoltz,H.
TITLE Plant inhibitors of fungal polygalacturonases and their use to control fungal disease
JOURNAL Patent: US 5569830-A 5 29-OCT-1996;
FEATURES Location/Qualifiers
source 1..215
/organism="unknown"
BASE COUNT 15 a 8 c 25 g 26 t 141 others
ORIGIN
Query Match 15.0%; Score 27; DB 25; Length 215;
Best Local Similarity 20.0%; Pred. No. 9.06e-03;
Matches 22; Conservative 42; Mismatches 45; Indels 1; Gaps 1;
Db 18 CNDKXKNGNTSSWTTDCNRTNGVCTDTTFRVNNDSGHMKYSANWYGGNNYGAAK 77
OY 58 CCTTGCGAGCTGACATGGAACAGGCGGGATGACTTTGCACTGAGAGCTG-AAAGAGT 116
Db 78 THYHTVNSGADSKTYTDSYNASCTSSNGTIDGNRSGADSYGSSKTAM 127
OY 117 CTTTCTGACAGTTCCTCTCATGAGTCGACGCTTCTGGAATTCCTGAA 166
RESULT 6 128278 215 bp DNA PAT 30-OCT-1996
LOCUS Sequence 5 from Patent US 5569830.
DEFINITION 128278
ACCESSION 128278
MID 91819054
VERSION 128278.1 GI:1819054
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 215)
AUTHORS Bennett,A., Labavitch,J.M., Powell,A. and Stoltz,H.
TITLE Plant inhibitors of fungal polygalacturonases and their use to control fungal disease
JOURNAL Patent: US 5569830-A 5 29-OCT-1996;
FEATURES Location/Qualifiers
source 1..215
/organism="unknown"
BASE COUNT 15 a 8 c 25 g 26 t 141 others
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Best Local Similarity 13.8%; Pred. No. 9.06e-03;
Matches 22; Conservative 66; Mismatches 69; Indels 2; Gaps 2;
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CP 158 ATTCCAGAGAGCTGACATGAGAGAGACTTGCAGAAAGACTCTTCACTTCACT 100
Db 61 YSSANWYGGNNYGAANTHYTHTVNSGADSKTYTDSYNASCTSSNGTIDGNRSGAD 119
CP 99 TGCNAAGCTATACCGCGCCCTGTTCCATGTGAGCTGCAAGGAGGCTCAGAGAGAGACA 40
Db 120 YGSSKTAMTSRNRGCKTANNAVDNRMGDASVSGDPKTK 158
CP 39 AGGGCGAGCCAGACCCATAGTGGCACTGCGCTCTG 1
RESULT 7 AR024229 965 bp DNA PAT 04-DEC-1998
LOCUS Sequence 22 from Patent US 5795961.
DEFINITION AR024229
ACCESSION AR024229
MID 93977523
VERSION AR024229.1 GI:3977523
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
Unclassified.

REFERENCE 1 (bases 1 to 965)
AUTHORS Wallace,T.Paul, Harris,W.J., Carr,F.J., Old,L.J., Welt,S. and Kitamura,K.
TITLE Recombinant human anti-Lewis b antibodies
JOURNAL Patent: US 5795961-A 22 18-AUG-1998;
FEATURES Location/Qualifiers
source 1..965
/organism="unknown"
BASE COUNT 192 a 170 c 226 g 205 t 172 others
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Best Local Similarity 20.0%; Pred. No. 3.54e-02;
Matches 18; Conservative 41; Mismatches 30; Indels 1; Gaps 1;
Db 781 GTRHTVNSGVASTSTCTASDYTTSMGVNRGRGMDYGGGTYTNGRGRVTVADTSS 940
CP 93 GTCATACCGCCCTGT-TCCATGTGAGCTGCCAGAGGCTCAGAGGAGAGAGAGGG 35
Db 841 NSRSYTAADTANYCYVRGSDSDGDY 870
CP 34 CAGCCAGACCCCATAGTGCCACTGCGCT 5
RESULT 8 AC005369 74371 bp DNA PRI 01-AUG-1998
LOCUS Homo sapiens chromosome 5, BAC clone 119j3 (LBNL H175), complete
DEFINITION sequence.
ACCESSION AC005369
MID 9367505
VERSION AC005369.1 GI:3367505
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Euteria;
Primates; Catarrhini; Hominoidea; Homo.
REFERENCE 1 (bases 1 to 74371)
AUTHORS Kimmerly,W., Bondoc,M., Cheng,J., Connolly,K.S., Gunning,K.M., Kader,K., Miguel,T., Miller,C., Pitluck,S., Pollard,M., Rojeski,H., Subramanian,S. and Martin,C.H.
TITLE Sequencing of human chromosome 5
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 74371)
AUTHORS Rieke,D.O.
TITLE Large Scale Sequence Analysis and Annotation with the Sequence Comparison Analysis (SCAN) System
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 74371)
AUTHORS Kimmerly,W., Bondoc,M., Cheng,J., Connolly,K.S., Gunning,K.M., Davis,C.A., Kader,K., Miguel,T., Pitluck,S., Pollard,M., Rojeski,H., Subramanian,S. and Martin,C.H.
TITLE Direct Submission
JOURNAL Submitted (01-AUG-1998) Human Genome Center, DOE Joint Genome Institute, Lawrence Berkeley National Laboratory, MS 74-157, Berkeley, CA 94720, U.S.A.
COMMENT Sequence submitted by:
DOE Joint Genome Institute.
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/rpt_family="Alu"
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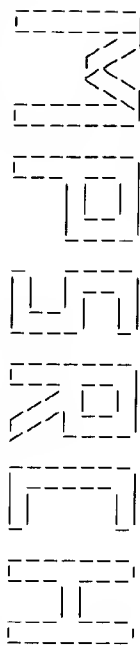
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                  6647..6684
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                  /rpt_type=tandem
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                  complement(7830..8185)
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                  9070..9387
repeat_region     /rpt_family="Alu"
                  complement(9740..9845)
repeat_region     /rpt_family="MER42"
                  complement(10440..11015)
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                  11950..12250
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                  12057..12085
                  /note="(A)29"
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                  /rpt_unit=A
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                  /rpt_unit=AC
                  13783..14024
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                  14175..14470
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                  /rpt_unit=GTTT
                  complement(19943..20222)
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                  complement(25727..26471)
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                  27774..28057
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                  28040..28066
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                  /rpt_unit=A
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                  28987..29214
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                  /note="958 identity dbSTS:G14522 (SHGC-11312)"
                  29495..29976
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                  /note="GRAIL 2 excellent exon, frame 2"
                  complement(30682..30733)
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                  complement(31573..31724)
                  /note="GRAIL 2 excellent exon, frame 1"
                  complement(32159..32232)
                  /note="GRAIL 2 excellent exon, frame 2"
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                  /db_xref="dbSTS:G26554"
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                  /note="GRAIL 2 excellent exon, frame 0"
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RESULT 11
LOCUS AC003030 60966 bp DNA PRI 06-JAN-1999
DEFINITION Homo sapiens chromosome 19, overlapping cosmids R29828 and F25496,
complete sequence.
ACCESSION AC003030
VERSION 94092821
KEYWORDS AC003030.1 GI:4092821
SOURCE HTG.
ORGANISM human.
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Primates; Catarrhini; Hominoidea; Homo.
REFERENCE
AUTHORS Lamerdin, J.E., McCready, P.M., Skowronski, E., Viswanathan, V.,
Burkhardt-Schultz, K., Gordon, L., Dias, J., Ramirez, M., Stillwagen, S.,
Phan, H., Velasco, N., Do, L., Regala, W., Terry, A., Gaines, J.,
Dangnan, L., Erler, A., Christensen, M., Georgescu, A., Avila, J.,
Liu, S., Attix, C., Andreise, T., Frankheim, M., Amico-Keller, G.,
Coeffield, J., Duarte, S., Lucas, S., Bruce, R., Thomas, P., Quan, G.,
Krommiller, B., Arellano, A., Sanders, C., Ow, D., Nolan, M., Truong, S.,
Kobayashi, A., Olsen, A.S. and Carrano, A.V.
TITLE Sequence analysis of a 2 Mb contig in 19p12 between UBA52 and
D19S455
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 60966)
AUTHORS Lamerdin, J.E.
TITLE Direct Submission
JOURNAL Submitted (28-OCT-1997) Human Genome Center, Joint Genome
Institute/ Lawrence Livermore National Laboratory, 7000 East Ave.,
Livermore, CA 94551, USA
REFERENCE 3 (bases 1 to 60966)
AUTHORS Lamerdin, J.E.
TITLE Direct Submission
JOURNAL Submitted (04-JAN-1999) Joint Genome Institute, Lawrence Livermore
National Laboratory, 7000 East Ave., Livermore, CA 94551, USA
REFERENCE 4 (bases 1 to 60966)
AUTHORS Lamerdin, J.E.
TITLE Direct Submission
JOURNAL Submitted (06-JAN-1999) Joint Genome Institute, Lawrence Livermore
National Laboratory, 7000 East Ave., Livermore, CA 94551, USA
COMMENT Map and sequence oriented from p telomere to centromere. This
accession is comprised of overlapping cosmids R29828 (bases 1 to
40,974) and F25496 (bases 23,336 to 60,966). R29828 is separated
from cosmid F23858 (NC004475) to the left by a sequence gap of
approximately 14 kb, which is to be filled by sequencing a
restriction fragment from cosmid R27236 (currently in progress).
Cosmid F25496 is separated from cosmid R31863 to the right by an
expected sequence gap of at least 10 kb. Additional chr 19 map and
sequence information are available at:
<http://www.bio.lnl.gov/genome/genome.html>.
LOCATION/Qualifiers
source
1. 60966
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="R29828-F25496"
/chromosome="19"
/map="19p12 between UBA52 and D19S455."
/cell_line="UV5HL9-5B for F25496, and 5HL2-B for R29828"
/clone_lib="IL19NC02 (for F25496) and IL19NC03 (for
R29828) chromosome 19-specific cosmid libraries"
/note="Cosmid libraries constructed at LBN from
flow-sorted chromosomes from human-hamster hybrid cell
lines UV5HL9-5B and 5HL2-B, each of which carries
chromosome 19 as its only human chromosome."
18. 219
/rpt_family="AluDo"
530. 824
repeat_region
/rpt_family="AluY"
930. 1066
/rpt_family="FLAM.C"
complement(1148..1315)
/note="predicted exon, program: graal2exons_human_1.3,
misc_feature

frame: 2, quality: good, score: 51.000"
complement(1477..1706)
/rpt_family="AluDo"
2105. 2406
repeat_region
/rpt_family="AluSp"
2513. 2809
repeat_region
/rpt_family="AluDo"
5028. 5045
misc_feature
/note="DGS similarity to overlapping ESTs: AA963316
UI-R-El-g1-c-06-0-UI-s1 UI-R-El Rattus norvegicus CDNA
clone UI-R-El-g1-c-06-0-UI-3' (220..237); 1008
identity. AA993275 ES197078 Normalized rat kidney, Bento
Scores Rattus sp. CDNA clone RKB38 3' end; (240..257);
1008 identity.
join(5028..5045,5432..5497,6480..6575,8155..8249,
10156..10183,10562..10717,11845..11991,12086..12176,
13147..13250,14614..14679,18159..18248,18465..18542,
18635..18746,23665..23755,24574..24709,37283..37493)
/note="Hypothetical human protein (partial)"
/codon_start=1
/evidence=not_experimental
/product="R29828.1"
/protein_id="AA03161.1"
/db_xref="PIR:94106983"
/db_xref="GI:4106983"
/translation="MISQIQDFEDVNFENASILSELTCQNSVDAAKPLKRAIOI
SOOTPYMCRLLFOLAOHLERDVSACDILGAGYARVVSYYTALFLSKML
LMERLQEVHPLLTGLQIVENMOGNFIOESRPFYLYVLTATYADNOVSVPK
IKLOQCIOTISTJHDEILPSPADLPFHMPEKHCQVLYVTVMHKNQGLIERAO
KYDRAIMQLEKLMKDCSPILSSFOVILHIIIMKRLTYGKATKADQGLICVNC
MDNAEOPFTALRLYSILIRINPDSFVSSCLRNARFTVGLSFPGIEVNAF
LPRULKMSARDIRLACGLVGLHITVIGNRHSNMVVPAMQAKIKPMQVQI
WSSALRLQKACGNADNHEAQNQHFQSOQLQDHIACSLPQHNLITVFGHMG
LAAIRLPVAFMEGSGSTTLYSADIKWKLQEDTESSSAWAPFRCIKSLPOTRAII
LLEPDL"
complement(5186..5264)
/note="Predicted exon, program: graal2exons_human_1.3,
frame: 0, quality: excellent, score: 86.000"
5432. 5497
misc_feature
/note="DGS similarity to overlapping ESTs: AA963316
UI-R-El-g1-c-06-0-UI-s1 UI-R-El Rattus norvegicus CDNA
clone UI-R-El-g1-c-06-0-UI-3' (238..303); 88%
identity. AA993275 ES197078 Normalized rat kidney, Bento
Scores Rattus sp. CDNA clone RKB38 3' end; (258..323);
88% identity.
complement(5602..5883)
/rpt_family="AluDo"
6480. 6575
repeat_region
/note="DGS similarity to overlapping ESTs: AA963316
UI-R-El-g1-c-06-0-UI-s1 UI-R-El Rattus norvegicus CDNA
clone UI-R-El-g1-c-06-0-UI-3' (304..399); 89%
identity. AA993275 ES197078 Normalized rat kidney, Bento
Scores Rattus sp. CDNA clone RKB38 3' end; (324..419);
89% identity. (6480..6660) predicted exon, program:
graal2exons_human_1.3, frame: 2, quality: excellent,
score: 76.000"
8155. 8249
misc_feature
/note="predicted exon, program: graal2exons_human_1.3,
frame: 0, quality: excellent, score: 94.000--DGS
similarity to overlapping ESTs: (8155..8237) AA963316
UI-R-El-g1-c-06-0-UI-s1 UI-R-El Rattus norvegicus CDNA
clone UI-R-El-g1-c-06-0-UI-3' (400..482); 88%
identity. (8155..8240) AA993275 ES197078 Normalized rat
kidney, Bento Scores Rattus sp. CDNA clone RKB38 3' end;
(420..505); 85% identity."
8351. 8477
repeat_region
/rpt_family="AluUb"
8480. 8768
repeat_region
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8778. 8837
misc_feature
/note="predicted exon, program: graal2exons_human_1.3,
frame: 2, quality: excellent, score: 83.000"
complement(8872..8996)
repeat_region

KEYWORDS	unidentified.
SOURCE	unidentified.
ORGANISM	unclassified.
REFERENCE	1 (bases 1 to 60)
AUTHORS	Oerum,H. and Seeger,C.
TITLE	METHOD FOR GENERATING MULTIPLE DOUBLE STRANDED NUCLEIC ACIDS
JOURNAL	Patent: WO 9720068-A 1 05-JUN-1997;
FEATURES	BOHRINGER MANHEIM GMBH (DE)
SOURCE	Location/Qualifiers
	1..60
	/organism="unidentified"
	/db_xref="taxon:32644"
BASE COUNT	0 a 30 c 0 g 0 t 30 others
ORIGIN	
Query Match	13.3%; Score 24; DB 25; Length 60;
Best Local Similarity	6.2%; Pred. No.5.03e-01;
Matches	2; Conservative 26; Mismatches 4; Indels 0; Gaps 0;
Db	29 CCRRRRRRRRRRRRRRRRRRRRRRRRRRRR 60
Cp	63 CCACGAGGCTCAAGAGGAGACACAGGCGCAC 32
RESULT 13	
LOCUS	MVU87256 1056 bp DNA MAM 02-JAN-1999
DEFINITION	Mustela vison GR dinucleotide repeat, chromosome 1q.
ACCESSION	U87256
NID	94099442
VERSION	U87256.1 GI:4099442
KEYWORDS	American mink.
SOURCE	Mustela vison
ORGANISM	Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
	Eumariotae; Fissipedia; Mustellidae; Mustela.
REFERENCE	1 (bases 1 to 1056)
AUTHORS	Bursgaard,K., Shukr,L.N.M., Malchenko,S., Koroleva,I. and Lohi,O.
TITLE	Direct Submision
JOURNAL	Submitted (27-JAN-1997) Breeding and Genetics, Danish Institute of
	Animal Science, Blichersalle K25, Tjele 8830, DK
FEATURES	Location/Qualifiers
source	1..1056
	/organism="Mustela vison"
	/db_xref="taxon:9667"
	/chromosome="1"
	/map="1q"
	/note="Primers: 1167F: agccctgcatactactctt, 1167R:
	gagactctaccgcctgttgag
	98..119
	/standard_name="1167F"
	complement(300..320)
	/standard_name="1167R"
BASE COUNT	211 a 221 c 210 g 225 t 189 others
ORIGIN	
Query Match	13.3%; Score 24; DB 23; Length 1056;
Best Local Similarity	10.8%; Pred. No.5.03e-01;
Matches	7; Conservative 40; Mismatches 17; Indels 1; Gaps 1;
Db	611 KRYRRCRYMMARVCVGSCTVARRCCCDKSGSCDKSHSKRYKMYMDRYHBC-KSMCA 669
Cp	99 TCGAAGTCATACACCGCGCCCTTTCATGTGAGCGTCCACAGAGCGTCACAGAGCAGACA 40
Db	670 MYNCK 674
Cp	39 AGGCG 35
RESULT 14	
LOCUS	MVU87256 1056 bp DNA MAM 02-JAN-1999
DEFINITION	Mustela vison GR dinucleotide repeat, chromosome 1q.
ACCESSION	U87256



(TM)

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MPerch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 17:32:37 1999; MasPar time 60.50 Seconds

Tabular output not generated. 637,452 Million cell updates/sec

Title: >US-09-092-296-3
Description: (1-180) from US09092296.seq
Perfect Score: 180
N.A. sequence: 1 CAGGAGCGAGTGGCCACTA.....CTTGAAGCTCTGCCCT 180
Comp: GTCTCGCGTCCACGGTGAT.....GAACCTTCGAGACGAGGA

Scoring table: TABLE default

Gap 6

Mmatch STD : Dbase 0; Query 0

Searched: 271905 seqs, 107135622 bases x 2

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database:

n-geneseg35
1:part1 2:part2 3:part3 4:part4 5:part5 6:part6 7:part7
8:part8 9:part9 10:part10 11:part11 12:part12 13:part13
14:part14 15:part15 16:part16 17:part17 18:part18
19:part19 20:part20 21:part21 22:part22 23:part23
24:part24 25:part25 26:part26 27:part27 28:part28
29:part29 30:part30 31:part31 32:part32 33:part33
34:part34 35:part35 36:part36 37:part37 38:part38
39:part39 40:part40 41:part41 42:part42 43:part43
44:part44 45:part45 46:part46 47:part47 48:part48
49:part49 50:part50 51:part51 52:part52 53:part53
54:part54 55:part55 56:part56 57:part57 58:part58
59:part59 60:part60

Statistics: Mean 7.535; Variance 4.714; scale 1.598

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description	Pred. No.
1	178	98.9	439	60	V84366 Human stomach carcinoma	4.41e-99
2	37	20.6	91	9	O51746 Oligonucleotide probe	1.15e-08
3	37	20.6	204	1	N81164 Base substituted E.co	1.15e-08
4	34	18.9	91	9	O51746 Oligonucleotide probe	5.55e-07
5	33	18.3	204	1	N81164 Base substituted E.co	1.99e-06
6	31	17.2	114	12	070467 Generic DNA sequence	8.61e-05
7	30	16.7	114	12	070470 Generic DNA sequence	2.96e-04
8	29	16.1	114	12	070470 Generic DNA sequence	2.96e-04
9	29	16.1	114	12	070467 Generic DNA sequence	2.96e-04

10	28	15.6	114	12	070469 Generic DNA sequence	1.00e-03
11	28	15.6	114	12	070465 Generic DNA sequence	1.00e-03
12	28	15.6	114	12	070468 Generic DNA sequence	1.00e-03
13	28	15.6	178	32	T76405 Human endothelin-1 an	1.00e-03
14	27	15.0	114	12	070468 Generic DNA sequence	3.36e-03
15	27	15.0	114	12	070465 Generic DNA sequence	3.36e-03
16	26	14.4	114	12	070472 Generic DNA sequence	1.11e-02
17	26	14.4	114	12	070466 Generic DNA sequence	1.11e-02
18	25	13.9	39	7	O51787 Mixed oligonucleotide	3.60e-02
19	25	13.9	114	12	070469 Generic DNA sequence	1.15e-01
20	24	13.3	114	12	070472 Generic DNA sequence	1.15e-01
21	24	13.3	114	12	070473 Generic DNA sequence	3.62e-01
22	23	12.8	114	12	070471 Generic DNA sequence	3.62e-01
23	23	12.8	114	12	070473 Generic DNA sequence	3.62e-01
24	23	12.8	172	32	T76363 HCV envelope region n	3.62e-01
25	23	12.8	565	6	O35072 Ballast Constituent c	1.11e+00
26	22	12.2	36	2	O11195 DC43 TSAR library gen	1.11e+00
27	22	12.2	75	21	T13612 DC43 TSAR library gen	1.11e+00
28	22	12.2	82	21	T13610 Human IL5 antisense o	1.11e+00
29	22	12.2	89	32	T76219 Generic DNA sequence	1.11e+00
30	22	12.2	114	12	070471 Generic DNA sequence	1.11e+00
31	22	12.2	114	12	070473 Substance P antisense	1.11e+00
32	22	12.2	250	32	T76438 Sequence encoding new	1.11e+00
33	22	12.2	264	32	T76445 Sequence encoding new	1.11e+00
34	22	12.2	501	3	N50025 ADP ribosylation fact	1.11e+00
35	22	12.2	501	3	N50026 TSAR-9 library genera	3.36e+00
36	22	12.2	3088	16	T13585 TSAR-9 library genera	3.36e+00
37	22	11.7	66	21	T13583 DC43 TSAR library gen	3.36e+00
38	21	11.7	69	21	T13583 B. malayi ankryrin cDN	3.36e+00
39	21	11.7	82	21	T13610 D. immitis ankryrin cD	3.36e+00
40	21	11.7	908	51	V63025 D. immitis ankryrin nd	3.36e+00
41	21	11.7	908	51	V63024 D. immitis ankryrin cd	3.36e+00
42	21	11.7	909	51	V63015 D. immitis ankryrin cd	3.36e+00
43	21	11.7	909	51	V63014 D. immitis ankryrin cd	3.36e+00
44	21	11.7	5235	51	V63023 D. immitis ankryrin cd	3.36e+00
45	21	11.7	5503	51	V63021 D. immitis ankryrin cd	3.36e+00

ALIGNMENTS

RESULT 1
ID V84366 standard; cDNA to mRNA; 439 BP.
AC V84366;
DT 30-MAR-1999 (first entry)
DE Human stomach carcinoma cDNA clone HP10408.
KW Transmembrane protein; HP10408; human; stomach cancer; ds.
OS Homo sapiens.
FH Key Location/Qualifiers
FT 75..311
FT /*tag a
FT /*note- "cDNA comprising the coding region (minus
the stop codon) is claimed (Claim 3)"
FN W09855508-f2.
PD 10-DEC-1998.
PF 03-JUN-1998; J02445.
PR 03-JUN-1997; JP-144948.
PA (PROT-) PROTEGENE INC.
PI (SAGA) SAGAMI CHEM RES CENTRE.
PI Kato S, Sekine S, Yamaguchi T;
WP: 99-045730/04.
P-PSDB; W08498.
PT New human proteins containing transmembrane domains and their
encoding sequences - useful in the preparation of antibodies and
large-scale protein production, gene diagnosis, and gene therapy
PS Claim 4; Page 135; 178pp; English.
CC This is the nucleotide sequence of cDNA clone HP10408, which
includes a coding region (also claimed) for a novel human
transmembrane protein (see W08498). The clone was isolated from a
stomach cancer cDNA library using a signal sequence detection
method, and by protein synthesis by in vitro translation. The
encoded protein has a putative signal sequence and a putative
internal transmembrane domain. The invention provides nucleotide
sequences (see V84359-76) coding for 18 transmembrane proteins

```

FT Primer_bind 18/..204
PN EP-285123-A. /*tag= b
PD 05-MAY-1988.
PR 03-MAR-1988; 105163.
PR 03-APR-1987; US-034819.
PA (SUSO) SIONEM SOKERI OY.
PI Lehtovaara P., Knowles J., Koivula A., Bamford J., Reinikainen T.;
DR WPI; 88-279927/40.
FT Introducing random point mutations into nucleic acids -
FT by prepn of single stranded template, annealing a primer, elongation,
FT misincorporation, completion of molecules and screening.
PS Disclosure; P; English.
CC Random point mutations were introduced into the alpha fragment of
CC E.coli beta-galactosidase. The wild type sequence was obtained as a
CC single stranded template and an oligonucleotide was hybridised to
CC it to generate a popn of DNA molecules which terminate at all
CC possible nucleotide positions within a specified region. The
CC variable 3' ends generated in this way are used as primers for
CC reverse transcriptase. Nucleotides are misincorporated by the
CC transcriptase and the molecules are completed to forms that can be
CC amplified and then expressed in a suitable host-vector system.
CC The sequence covers all 1/6 given base substitutions, most of which
CC occurred singularly in any given mutant.
SQ See also P80575.
SQ Sequence 204 BP; 21 A; 47 C; 17 G; 11 T; 108 Others;
Query Match 20.6%; Score 37; DB 1; Length 204;
Best Local Similarity 7.9%; Pred. No. 1,15e-08;
Matches 8; Conservative 54; Mismatches 39; Indels 0; Gaps 0;
Db 86 yrrtthhyrrmbhnyridyngdsaaayccyrsvkydcynachdhhyvbybbyrnv 145
Cp 141 TCTAGAGAGGACTCTGCAGAAAGACCTCTCAGTTCAGTGCAGACGACATACCGCGC 82
Db 146 hmhmnccebnhvcnhybhnbnhwayrtharddvhc 186
Cp 81 CCTGTTCATGTGAGCTGCAGGAGGAGGTCAAGAGAGGAC 41
RESULT 4
ID 051746 standard; cDNA; 91 BP.
AC 051745:
DT 31-MAY-1994 (first entry)
DE Oligonucleotide probe MK14-A
OI Oligonucleotide; DNA probe; mycobacteria; disease diagnosis;
ss.
OS Synthetic.
PN EP-571911-A.
PD 01-DEC-1993.
PF 24-MAY-1993; 108325.
PR 26-MAY-1992; US-889651.
PA (BECT ) BECTION DICKINSON CO.
PI Shank DP, Spears PA;
PT WPI; 93-378844/48
DR New 0199:nucleotide probes specific for Mycobacteria - used for
FT detection and amplification of Mycobacteria nucleic acid in
FT samples
PS Claim 3; Page 14; 23pp; English.
CC Oligonucleotide probe MK14-A consists of nucleotides 5-95 of MK14
CC (051735). It hybridized to all spp. of mycobacteria tested, but
CC cross reacted to a few non-mycobacterial spp. The probe may
CC be useful as an initial screen for mycobacterial infection.
CC See also 051735-45 and 051747-59.
SQ Sequence 91 BP; 5 A; 17 C; 15 G; 4 T;
Query Match 18.9%; Score 34; DB 9; Length 91;
Best Local Similarity 10.7%; Pred. No. 5.55e-07;
Matches 6; Conservative 39; Mismatches 11; Indels 0; Gaps 0
Db 12 svhyvvvhhvshhsvvvvhhvsvvvvhhvvhvhhvhyvsvctcaagc 67
52 GACCTCTTGACACATGCAGACGAGCGCGGATATGACTTTCGACACTGAGAC 107

```

```

ID AC N81164 standard; DNA; 204 BP.
DE DT 08-NOV-1990 (first entry)
KW DE Base substituted E.coli beta-galactosidase alpha-fragment.
OS E.coli beta galactosidase alpha-fragment; base substitutions; ss.
FH Key Escherichia coli.
FT misc_feature Location/Qualifiers
FT primer_bind //tag= a
FT primer_bind //function=multiple cloning site
FT primer_bind 187..204
FT //tag= b
PN EP-285123-A.
PN 05-MAY-1988.
PN 30-MAR-1988; 105163.
PR 03-APR-1987; US-034819.
PA (SUZO) SUOMEN SOKERI OY.
PI Laitovaara P., Knowles J., Koivu A., Bamford J., Reinikainen T.;
PT WPI: 88-279927/40.
PT Introducing random point mutations into nucleic acids -
PT by prep of single stranded template, annealing a primer, elongation,
PT misincorporation, completion of molecules and screening.
PS Disclosure; P: English.
CC Random point mutations were introduced into the alpha fragment of
CC E.coli beta-galactosidase. The wild type sequence was obtained as a
CC single stranded template and an oligonucleotide was hybridised to
CC it to generate a popn of DNA molecules which terminate at all
CC possible nucleotide positions within a specified region. The
CC variable 3' ends generated in this way are used as primers for
CC reverse transcriptase. Nucleotides are misincorporated by the
CC transcriptase and the molecules are completed to forms that can be
CC amplified and then expressed in a suitable host-vector system.
CC The sequence covers all 176 diff base substitutions, most of which
CC occurred singularly in any given mutant.
SQ See also P80575.
SQ Sequence 204 BP; 21 A; 47 C; 17 G; 11 T; 108 Others;
Query Match 18.3%; Score 33; DB 1; Length 204;
Best Local Similarity 5.4%; Pred. No. 1.99e-06;
Matches 5; Conservative 51; Mismatches 37; Indels 0; Gaps 0;
Db 94 yirmdbnydyngnsdaawycgcrsykygcpcnaehdhyrybbbyrvnhmnc 153
OY 1 cagagaccatggccacmrtatggctgtggcgctgcccttccccctttgacctcct 60
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
Db 154 ccbanhcvbnvhnbnrhayvrhdarrdhvc 186
OY 61 tggcagctcatgatgaacagcgccgctatgac 93
RESULT
ID ID Q70467 standard; DNA; 114 BP.
AC AC Q70467.
DT DT 05-APR-1995 (first entry)
DE DE Generic DNA sequence to generate a random TSAR peptide library.
KW KW TSAR: totally synthetic affinity reagent; synthetic; binding domain;
KW effector domain; concatenated heterofunctional protein; linker;
KW direct; rapid; detection; screening; treatment; generati; ss.
OS OS Synthetic.
FH FH key Location/Qualifiers
FH key 55..60
FT misc_feature //tag= a
FT //note= "this sequence represents 'Z'; Z can be a
FT sequence of 6, 9 or 12 nucleotides (see
FT comments)"
PN PN MO9418318-A.
PN PD 18-AUG-1984.
PF PF 01-FEB-1994; U00977.
PF PF 01-FEB-1993; US-013416.
PR PR 30-DEC-1993; US-176500.

```

[illegible]

Query Match	16.78;	Score 30;	DB 12;	Length 114;
Best Local Similarity	6.78;	Pred. No. 8.61e-05;		
Matches	7;	Conservative	28;	Mismatches 69;
			Indels	0;
			Gaps	0;

RESULT	8
ID	Q70470 standard; DNA; 114 BP.

DT 10-Apr-1995 (first entry)
DE Generic DNA sequence to generate a random TSAR peptide library.
KW TSAR, totally synthetic affinity reagent; synthetic; binding domain
KW effector domain; concatenated heterofunctional protein; linker
KW direct; rapid; detection; screening; treatment; generic; ss.

EH	Key	Location/Qualifiers
FT	misc_feature	55..60
FT		/*tag= a
FT		/note="encoded by Z (see comments)"

```

FT      /note="encoded by z (see comments)"
PN      M09418318-A.
PD      18-AUG-1994.
PE      01-FEB-1994.    U00927.
PR      01-FEB-1993;    US-013416.
PR      30-DEC-1993;    US-176500.
PR      31-JAN-1994;    US-189531.
PA      (UNCL-) UNIT NORTH CAROLINA.
PI      Fowlkes DM, Kay BK.
DR      WPI; 94-27939/34.
        P-PSDB; R58378.

```

PT Identifying proteins or peptide(s) which bind a ligand - by
 PT screening a recombinant vector library expressing fusion proteins
 PT comprising a binding domain and an effector domain
 PS Disclosure: Page 35: 255pp: English.
 Q70470 is a generic DNA sequence used to generate random TSAR (Totally
 CC Synthetic Affinity Reagents) peptides. This generic formula can also be
 CC represented as follows: X(NNN)4(CAC)(NNN)4(CAC)(NNN)6(CAC)(NNN)8
 CC -(CAC)2(NNN)X. X and Y are flanking restriction sites (X is not the same
 CC as Y) that are not specified further. The peptides generated by this and
 CC other generic sequences (Q70471-73) have invariant histidine residues
 CC incorporated into variant sequences. TSARs are concatenated
 CC heterofunctional proteins or peptides, comprising at least two functional
 CC regions - a binding domain with affinity for a ligand and a second
 CC effector peptide portion that is chemically or biologically active. They
 CC may further comprise a linker peptide between the 2 domains. The TSARs
 CC or comps. comprising a TSAR binding domain can be used in vivo to

CC deliver chemically or biologically active moiety, eg. metal ion, the
CC radioisotope, peptide, toxin or enzyme, to the specific target or on the
CC cell. They can also replace the function of macromolecules, eg.
CC monoclonal or polyclonal antibodies and therefore circumvent the need
CC for complex methods of hybridoma formation or in vivo antibody
CC production. The TSARs are easily characterised and have designed
CC activity allowing direct and rapid detection in a screening process.
Sequence 114 bp; 5' A; 10 C; 0 G; 0 T.

Query Match	16.1%;	Score 29;	DB 12;	Length 114;
Best Local Similarity	7.5%;	Pred. No. 2.96e-04;		
Matches	7;	Conservative	25;	Mismatches 61;
			Indels	0;
			Gaps	0;

[illegible]

RESULT	9
ID	Q70467 standard; DNA; 114 BP.
	Q70467

DI	05-APR-1995	(first entry)
DE	Generic DNA sequence to generate a random TSAR peptide library.	
KW	TSAR: totally synthetic affinity reagent; synthetic; binding domain	
KW	effector domain; concatenated heterofunctional protein; linker	
KW	direct; rapid; detection; screening; treatment; generic; ss.	
OS	Synthetic.	
PH	Key	Location/Qualifiers
FI	misc-feature	55..60

FT /note= "this sequence represents 'Z'; Z can be a
FT sequence of 6, 9 or 12 nucleotides (see
FT comments)"

PN MO9418318-A.
PD 18-AUG-1994.
PE 01-FEB-1994; U00977.
PR 01-FEB-1993; US-03416.
PR 30-DEC-1993; US-176500.
PR 31-JAN-1994; US-189331.
PA (UYNC-) UNIV NORTH CAROLINA.
PT Fovlkes DM, Kay BK;
DR WPI; 94-279739/34.
P-PSDB: R65153.

identifying proteins or peptide(s) which bind a ligand - by
 screening a recombinant vector library expressing fusion proteins
 comprising a binding domain and an effector domain
 Disclosing: Page 35; 255pp; English.
 Q70467 is a generic DNA sequence used to generate random TSAR (Totally
 Synthetic Affinity Reagents) peptides. This generic formula can be
 represented as follows: X(NNH)₁₆(TGC)(NNB)₁₂(NNB)₁₆(TGC)(NNB)₁₆. X
 and Y are flanking restriction sites (X is not the same as Y) that are
 not specified further. Other generic sequences are shown in Q70466-68.
 Other specific peptides generated by these generic sequences are shown in
 R5131-54. TSARs are concatenated heterofunctional proteins or peptides,
 comprising at least two functional regions - a binding domain with
 affinity for a ligand and a second effector peptide portion that is
 chemically or biologically active. They may further comprise a linker
 peptide between the 2 domains. The oligonucleotides are also designed so
 that the expressed peptide contains 2 or 4 cysteine residues positioned
 in, or flanking, the unpredicted or variant residues. These residues
 confer some degree of conformational rigidity to the peptides. The TSARs
 or compenss, comprising a TSAR binding domain can be used in vivo to
 deliver a chemically or biologically active moiety, eg. metal ion,
 radioisotope, peptide, toxin or enzyme, to the specific target or on the
 cell. They can also replace the function of macromolecules, eg.
 monoclonal or polyclonal antibodies and therefore circumvent the need for
 complex methods of hybridoma formation or in vivo antibody production.
 The TSARs are easily characterised and have designed actively allowing
 direct and rapid detection in a screening process.


```

RESULT 12
ID Q70468 standard; DNA; 114 BP.
AC Q70468;
DE 05-APR-1995 (first entry)
DE Generic DNA sequence to generate a random TSAR peptide library.
KW TSAR: totally synthetic affinity reagent; synthetic; binding domain;
KW effector domain; concatenated heterofunctional protein; linker;
KW direct; rapid; detection; screening; treatment; generic; ss.
OS Synthetic.
FH Key
FT misc_feature
FT 55..60 Location/Qualifiers
FT /*tag= a
FT /note= "this sequence represents '2'; 2 can be a
FT sequence of 6, 9 or 12 nucleotides (see
FT comments)"
PN W09418318-A.
PD 18-AUG-1994.
PF 01-FEB-1994; U00977.
PR 01-FEB-1993; US-013416.
PR 30-DEC-1993; US-176500.
PR 31-JAN-1994; US-189331.
PA (UYNC-) UNIV NORTH CAROLINA.
PI Fowlkes DM. Key BK.
PI WPI: 94-279739/34.
DR P-PSDB; R65134.
PT Identifying proteins or peptide(s) which bind a ligand - by
PT screening a recombinant vector library expressing fusion proteins
PT comprising a binding domain and an effector domain
PS Disclosure: Page 35; 255pp; English.
CC Q70468 is a generic DNA sequence used to generate random TSAR (Totally
CC Synthetic Affinity Reagents) peptides. This generic formula can also be
CC represented as follows: X(NNB)11(TGC)(NNB)7(TGC)(NNB)10Y. X
CC and Y are flanking restriction sites (X is not the same as Y) that are
CC not specified further. Other generic sequences are shown in Q70466-68.
CC Other specific peptides generated by these generic sequences are shown in
CC R65151-54. TSARs are concatenated heterofunctional proteins or peptides,
CC comprising at least two functional regions - a binding domain with
CC affinity for a ligand and a second effector peptide portion that is
CC chemically or biologically active. They may further comprise a linker
CC peptide between the 2 domains. The oligonucleotides are also designed so
CC that the expressed peptide contains 2 or 4 cysteine residues positioned
CC in, or flanking, the unpredicted or variant residues. These residues
CC confer some degree of conformational rigidity to the peptides. The TSARs
CC or compens. comprising a TSAR binding domain can be used in vivo to
CC deliver a chemically or biologically active moiety, eg. metal ion,
CC radioisotope, peptide, toxin or enzyme, to the specific target or on the
CC cell. They can also replace the function of macromolecules, eg.
CC monoclonal or polyclonal antibodies and therefore circumvent the need
CC for complex methods of hybridoma formation or in vivo antibody
CC production. The TSARs are easily characterised and have designed activity
CC allowing direct and rapid detection in a screening process.
CC Sequence 114 BP: 0 A; 2 C; 2 G; 2 T;
SQ
Query Match 15.6%; Score 28; DB 12; Length 114;
Best Local Similarity 2.8%; Pred. No. 1,00e-03;
Matches 3; Conservative 30; Mismatches 73; Indels 0; Gaps 0;

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```

KW Chronic obstructive pulmonary disease; bronchitis; ss.
OS Synthetic.
PN W09640162-A1.
PD 19-DEC-1996.
PF 06-JUN-1996; U09306.
PR 07-JUN-1995; US-474497.
PA (UYEC-) UNIV EAST CAROLINA.
PI Metzger WJ, NYCE JW.
PI WPI: 97-051871/05.
DR Treatment of airway diseases such as asthma - by topically applying
DR adenosine-free antisense oligo:nucleotide to airway epithelium of
PT subject
PT Claim 5; Page 38; 71pp; English.
CC A method for treating airway disease in a subject has been produced,
CC which involves the topical administration of an essentially adenosine
CC free antisense oligonucleotide (ON) to the airway epithelium of the
CC subject. The present sequence is an antisense oligonucleotide specific
CC for the human endothelin-1, targeted at the initiation codon. The
CC method can be used to treat airway diseases such as cystic fibrosis,
CC asthma, chronic obstructive pulmonary disease, bronchitis and other
CC airway diseases characterised by an inflammatory response. By
CC eliminating adenosine from the antisense ON, its liberation upon
CC antisense degradation is prevented, thereby preventing adenosine-
CC induced bronchoconstriction in patients with hyper-reactive airways.
CC Sequence 178 BP: 0 A; 52 C; 46 G; 32 T;
SQ
Query Match 15.6%; Score 28; DB 32; Length 178;
Best Local Similarity 25.0%; Pred. No. 1,00e-03;
Matches 18; Conservative 32; Mismatches 22; Indels 0; Gaps 0;

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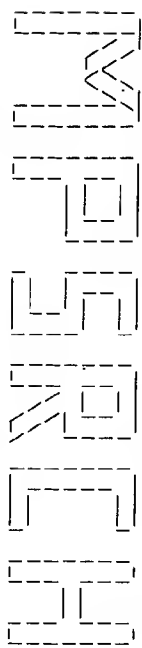
DB 100 cttggcggccgcbdbbcbgcbgbbdbbcbtcbgcbdbbcbtctgbbbbb 159
QY 27 cttggcggccgcbdbbcbgcbgbbdbbcbtcbgcbdbbcbtctgbbbbb 86
DB 160 ggggtcbbdbbc 171
QY 87 gttatgatttgc 98

```

```

RESULT 14
ID Q70468 standard; DNA; 114 BP.
AC Q70468;
DE 05-APR-1995 (first entry)
DE Generic DNA sequence to generate a random TSAR peptide library.
KW TSAR: totally synthetic affinity reagent; synthetic; binding domain;
KW effector domain; concatenated heterofunctional protein; linker;
KW direct; rapid; detection; screening; treatment; generic; ss.
OS Synthetic.
FH Key
FT misc_feature
FT 55..60 Location/Qualifiers
FT /*tag= a
FT /note= "this sequence represents '2'; 2 can be a
FT sequence of 6, 9 or 12 nucleotides (see
FT comments)"
PN W09418318-A.
PD 18-AUG-1994.
PF 01-FEB-1994; U00977.
PR 01-FEB-1993; US-013416.
PR 30-DEC-1993; US-176500.
PR 31-JAN-1994; US-189331.
PA (UYNC-) UNIV NORTH CAROLINA.
PI Fowlkes DM. Key BK.
PI WPI: 94-279739/34.
DR P-PSDB; R65154.
PT Identifying proteins or peptide(s) which bind a ligand - by
PT screening a recombinant vector library expressing fusion proteins
PT comprising a binding domain and an effector domain
PS Disclosure: Page 35; 255pp; English.
CC Q70468 is a generic DNA sequence used to generate random TSAR (Totally
CC Synthetic Affinity Reagents) peptides. This generic formula can also be
CC represented as follows: X(NNB)11(TGC)(NNB)6(TGC)(NNB)10Y. X
CC and Y are flanking restriction sites (X is not the same as Y) that are
CC not specified further. Other generic sequences are shown in Q70466-68.

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(TM)

Release 3.1a John F. Collins, Biocomputing Research Unit.
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Distribution rights by Oxford Molecular Ltd

Mperch_lm n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 17:33:58 1999; Master time 16.98 Seconds

Tabular output not generated. 916.867 Million cell updates/sec

Title: >US-09-092-296-3

Description: (1-180) from US09092296.seq

Perfect Score: 180 1 CAGGAGGCGAGTGGCCACCTA.....CTTGAAGCTGTCGCCCT 180

N.A. Sequence: GTCCTGCGGTGACCGGTGAT.....GAACCTTTGAGAGGAGAGA

Comp: 1 CAGGAGGCGAGTGGCCACCTA.....CTTGAAGCTGTCGCCCT 180

Scoring table: TABLE default

Gap 6

Mmatch STD: Dbase 0; Query 0

Searched: 165359 segs, 43243793 bases x 2

Post-processing: Minimum Match 08

Listing first 45 summaries

Database: n-issued

1:5A.COMB 2:5B.COMB 3:5C.COMB 4:PC79.COMB 5:backfiles1

Statistics: Mean 7.135; Variance 3.967; scale 1.799

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,

and is derived by analysis of the total score distribution.

SUMMARIES

Result

No. Score Match Length DB ID Description Pred. No.

1 29 16.1 965 3 US-08-388- Sequence 22, Applicat 6.29e-06

2 28 15.6 7218 2 US-08-232- Sequence 14, Applicat 2.53e-05

3 27 15.0 215 1 US-08-238- Sequence 5, Applicat 1.01e-04

4 26 15.0 215 1 US-08-238- Sequence 5, Applicat 1.01e-04

5 26 14.4 965 3 US-08-388- Sequence 22, Applicat 3.93e-04

6 22 12.2 82 4 PCT-US95-1 Sequence 99, Applicat 7.70e-02

7 22 12.2 3088 3 US-08-418- Sequence 97, Applicat 7.70e-02

8 22 12.2 3088 3 US-08-418- Sequence 97, Applicat 7.70e-02

21 21 11.7 906 3 US-08-847- Sequence 41, Applicat 2.73e-01

22 21 11.7 906 3 US-08-847- Sequence 40, Applicat 2.73e-01

23 21 11.7 906 3 US-09-031- Sequence 40, Applicat 2.73e-01

24 21 11.7 908 3 US-09-031- Sequence 39, Applicat 2.73e-01

25 21 11.7 908 3 US-08-847- Sequence 39, Applicat 2.73e-01

26 21 11.7 908 3 US-08-847- Sequence 37, Applicat 2.73e-01

27 21 11.7 908 3 US-09-031- Sequence 37, Applicat 2.73e-01

28 21 11.7 909 3 US-08-847- Sequence 26, Applicat 2.73e-01

29 21 11.7 909 3 US-09-031- Sequence 26, Applicat 2.73e-01

30 21 11.7 909 3 US-09-031- Sequence 25, Applicat 2.73e-01

31 21 11.7 909 3 US-08-847- Sequence 25, Applicat 2.73e-01

32 21 11.7 911 3 US-09-031- Sequence 24, Applicat 2.73e-01

33 21 11.7 911 3 US-08-847- Sequence 24, Applicat 2.73e-01

34 21 11.7 911 3 US-08-847- Sequence 22, Applicat 2.73e-01

35 21 11.7 911 3 US-09-031- Sequence 22, Applicat 2.73e-01

36 21 11.7 5235 3 US-09-031- Sequence 22, Applicat 2.73e-01

37 21 11.7 5235 3 US-08-847- Sequence 36, Applicat 2.73e-01

38 21 11.7 5235 3 US-09-031- Sequence 35, Applicat 2.73e-01

39 21 11.7 5235 3 US-08-847- Sequence 35, Applicat 2.73e-01

40 21 11.7 5503 3 US-09-031- Sequence 35, Applicat 2.73e-01

41 21 11.7 5503 3 US-08-847- Sequence 34, Applicat 2.73e-01

42 21 11.7 5503 3 US-08-847- Sequence 34, Applicat 2.73e-01

43 21 11.7 5503 3 US-09-031- Sequence 32, Applicat 2.73e-01

44 20 11.1 66 4 PCT-US95-1 Sequence 93, Applicat 5.46e-01

45 20 11.1 2334 1 US-08-406- Sequence 1, Applicat 9.46e-01

ALIGNMENTS

RESULT 1
ID US-08-388-672A-22 STANDARD; DNA; UNC; 965 BP.
AC xxxxxx

DE Sequence 22, Application US/08388672A
CC Sequence 22, Application US/08388672A
CC Patent No. 5793961

CC GENERAL INFORMATION:
CC APPLICANT: Wallace, T. Paul
CC APPLICANT: Harris, William J.

CC APPLICANT: Carr, Frank J.
CC APPLICANT: Old, Lloyd J.
CC APPLICANT: Welf, Sydney

CC APPLICANT: Kitamura, Kunio
CC TITLE OF INVENTION: Recombinant Human Anti-Lewis B
CC TITLE OF INVENTION: Antibodies
CC NUMBER OF SEQUENCES: 25

CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Felle and Lynch
CC STREET: 805 Third Avenue
CC CITY: New York
CC STATE: New York
CC COUNTRY: U.S.A.

CC ZIP: 10022
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible

CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patent Release #1.0, Version #1.30.
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/388.672A
CC FILING DATE: 14-FEB-1995

CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Hanson, No. 5795961man D.
CC REGISTRATION NUMBER: 30,946

CC REFERENCE/DOCKET NUMBER: LUD 5409
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: 212-688-9200
CC TELEFAX: 212-838-3864

CC INFORMATION FOR SEQ ID NO: 22:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 965 base pairs
CC TYPE: nucleic acid

CC STRANDEDNESS: unknown
CC TOPOLOGY: unknown
CC MOLECULE TYPE: DNA (genomic)
SQ SEQUENCE 965 BP; 192 A; 170 C; 226 G; 200 T; 177 OTHER.

Query Match 16.1%; Score 29; DB 3; Length 965;
Best Local Similarity 19.6%; Pred. No. 6,29e-06;
Matches 21; Conservative 48; Mismatches 37; Indels 1; Gaps 1;

Db 857 VGRGRTSDGDDYWGTTVTYVSSHVKDMTSSSASVGRVTCSSSTHNGNTYWK 916
Y 4 GACCCGAGTGGCAGCTATGGGGTCTGGGCTGCCCTTGCTCTGACC-CTCCTTG 62
Matches 21; Conservative 48; Mismatches 37; Indels 1; Gaps 1;

Db 917 GKAKYFVNSRSGSVSGSGDTYTSSDATYTCGTHARTGKYKG 963
Y 63 GCAGCTCAGATGGAACAGGCGCGGTATGACTTTCGACATGAAGCTG 109

RESULT 2
ID US-08-232-463-14 STANDARD; DNA; UNC; 7218 BP.
AC xxxxxx

DE Sequence 14, Application US/08232463
CC Sequence 14, Application US/08232463
CC Patent No. 5670367

GENERAL INFORMATION:
CC APPLICANT: DORNER, F.
CC APPLICANT: SCHEIFLINER, F.
CC APPLICANT: FALKNER, F. G.
CC TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUS
CC NUMBER OF SEQUENCES: 52
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Foley & Lardner
CC STREET: 1800 Diagonal Road, Suite 500
CC CITY: Alexandria
CC STATE: VA
CC COUNTRY: USA
CC ZIP: 22313-0299

COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/232,463
CC FILING DATE:
CC CLASSIFICATION: 435
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US/07/935,313
CC FILING DATE:
CC APPLICATION NUMBER: EP 91 114 300.6
CC FILING DATE: 26-AUG-1991

ATTORNEY/AGENT INFORMATION:
CC NAME: BENT, Stephen A.
CC REGISTRATION NUMBER: 29,768
CC REFERENCE/DOCKET NUMBER: 30472/114 IMNU
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (703)836-9300
CC TELEFAX: (703)683-4109
CC TELEX: 899149

INFORMATION FOR SEQ ID NO: 14:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 7218 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC IMMEDIATE SOURCE:
CC CLONE: pTZipt-Fls

SEQUENCE 7218 BP; 1944 A; 1491 C; 1486 G; 1929 T; 368 OTHER.

Query Match 15.6%; Score 28; DB 2; Length 7218;
Best Local Similarity 2.5%; Pred. No. 2,53e-05;
Matches 4; Conservative 90; Mismatches 66; Indels 0; Gaps 0;

Db 1061 TGGCATTYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYYY 1120
Y 21 TGGGCTGTGGCCCTGCTCTCTCTGACCTCTGGAGCTCAGTGAACAG 80

Db 1121 YY 1180
Y 81 GGCCGGGTATGACTTTCGACACTGAAGAGCTTTCTGCAAGTCTCTCATG 140

Db 1181 YY 1220
Y 141 AGTCAGCTTCTGGAATGCTGAAGAAAGCTCTGCTCTCT 180

RESULT 3
ID US-08-238-163-5 STANDARD; DNA; UNC; 215 BP.
AC xxxxxx

DE Sequence 5, Application US/08238163
CC Sequence 5, Application US/08238163
CC Patent No. 5569830

GENERAL INFORMATION:
CC APPLICANT: BENNETT, Alan
CC APPLICANT: LABAYTCH, John M.
CC APPLICANT: POWELL, Ann
CC APPLICANT: STOTZ, Henrik
CC TITLE OF INVENTION: PLANT INHIBITORS OF FUNGAL
CC NUMBER OF SEQUENCES: 24
CC CORESPONDENCE ADDRESS:
CC ADDRESSEE: Townsend and Townsend Kourie and Crew
CC STREET: Stewart Street Tower, One Market Plaza
CC CITY: San Francisco
CC STATE: California
CC COUNTRY: US
CC ZIP: 94105-1493

COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/238,163
CC FILING DATE: 03-MAY-1994
CC CLASSIFICATION: 800

ATTORNEY/AGENT INFORMATION:
CC NAME: Bastian, Kevin L.
CC REGISTRATION NUMBER: 34,774
CC REFERENCE/DOCKET NUMBER: 2307E-540
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (415) 543-9600
CC TELEFAX: (415) 543-5043

INFORMATION FOR SEQ ID NO: 5:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 215 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: unknown
CC MOLECULE TYPE: protein
CC FEATURE:
CC NAME/KEY: misc.feature
CC LOCATION: 1..215

OTHER INFORMATION: /standard_name="Deduced amino acid
CC OTHER INFORMATION: sequence of Peir from bean."
SQ SEQUENCE 215 BP; 15 A; 8 C; 25 G; 26 T; 141 OTHER.

Query Match 15.0%; Score 27; DB 1; Length 215;
Best Local Similarity 20.0%; Pred. No. 1,01e-04;
Matches 22; Conservative 42; Mismatches 45; Indels 1; Gaps 1;

Db 18 CNDKAKKGGNTTSSWTDCNRTWGVCDTITTYVNDSGHKKYSANYYGGNNYAAK 77
Y 58 CTTTGCAGCTCAGTGAACAGGCGCGGTATGACTTTCGCAACTGAAGCTG-AGGAGT 116


```

CC      TELEX: 248855 OPAT UR
CC      INFORMATION FOR SEQ ID NO: 1:
CC      SEQUENCE CHARACTERISTICS:
CC      LENGTH: 3088 base pairs
CC      TYPE: nucleic acid
CC      STRANDEDNESS: double
CC      TOPOLOGY: linear
CC      MOLECULE TYPE: DNA (genomic)
SQ      SEQUENCE 3088 BP; 716 A; 761 C; 672 G; 939 T; 0 OTHER.

Dn      Query Match          12.2%; score 22; DB 3; length 3088;
Dn      Best Local Similarity 72.0%; Pred. No. 7,70e+02;
Cc      Matches 36; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

Cc      15 CCGAAGATGGGCCAGAGGGGGGCCAGAGGGCTGCCAGGGGCGCCCTGTGGC 64
Cc      1111111111111111111111111111111111111111111111111111111
Cc      Cp      63 CCAAGAGGCTCAAGAGAGGAGACAGAGGGGGCCAGACCCCATATGTGGC 14

RESULT          9
ID      US-08-471-052A-145 STANDARD; DNA; UNC; 65 BP.
Dn      Axxxxx
Dn      Sequence 145, Application US/08471052A
Dn      Sequence 145, Application US/08471052A
Cc      Patent No. 5625033
Cc      GENERAL INFORMATION:
Cc      APPLICANT: Kay, B. K.
Cc      APPLICANT: Fowlkes, D. M.
Cc      TITLE OF INVENTION: Totally Synthetic Affinity Reagents
Cc      NUMBER OF SEQUENCES: 166
Cc      CORRESPONDENCE ADDRESS:
Cc      ADDRESSEE: Pennile & Edmonds
Cc      STREET: 1155 Avenue of the Americas
Cc      CITY: New York
Cc      STATE: New York
Cc      COUNTRY: U.S.A.
Cc      ZIP: 10036-2711
Cc      COMPUTER READABLE FORM:
Cc      MEDIUM TYPE: floppy disk
Cc      COMPUTER: IBM PC compatible
Cc      OPERATING SYSTEM: PC-DOS/MS-DOS
Cc      SOFTWARE: Patentin Release #1.0, Version #1.25
Cc      CURRENT APPLICATION DATA:
Cc      APPLICATION NUMBER: US/08/471,052A
Cc      FILING DATE: 06-JUNE-1995
Cc      CLASSIFICATION: 530
Cc      ATTORNEY/AGENT INFORMATION:
Cc      NAME: Mirock, S. Leslie
Cc      REGISTRATION NUMBER: 18,872
Cc      REFERENCE/DOCKET NUMBER: 1101-179
Cc      TELECOMMUNICATION INFORMATION:
Cc      TELEPHONE: 212 790-9090
Cc      TELEFAX: 212 869-8864/9741
Cc      TELEX: 66141 PENNILE
Cc      INFORMATION FOR SEQ ID NO: 145:
Cc      SEQUENCE CHARACTERISTICS:
Cc      LENGTH: 65 bases
Cc      TYPE: nucleic acid
Cc      STRANDEDNESS: single
Cc      TOPOLOGY: unknown
Cc      MOLECULE TYPE: DNA
Cc      SEQUENCE 65 BP; 3 A; 3 C; 3 G; 2 T; 54 OTHER.

Dn      Query Match          11.7%; score 21; DB 1; length 65;
Dn      Best Local Similarity 13.8%; Pred. No. 2,73e-01;
Cc      Matches 9; Conservative 16; Mismatches 40; Indels 0; Gaps 0;

Dn      1 CTAGAGNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNA 60
Dn      1111111111111111111111111111111111111111111111111111111
Cp      147 CTGAGCTCATAGAGGAGCACTTGTACAGAAAGACTCTTCACGCTTCAGTTGCAAGTGCAT 88
Dn      61 CCTGG 65

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Cc      87 CCGG 83          |||
                                |
RESULT 10                      |
ID     US-08-471-052A-144 STANDARD; DNA: UNC; 66 BP.
AC      xxxxxx
DE      Sequence 144, Application US/08471052A
CC      Sequence 144, Application US/08471052A
CC      Patent No. 5625033
CC      GENERAL INFORMATION:
CC      APPLICANT: Kay, B. K.
CC      APPLICANT: Fowles, D. M.
CC      TITLE OF INVENTION: Totally Synthetic Affinity Reagents
CC      NUMBER OF SEQUENCES: 166
CC      CORRESPONDENCE ADDRESS:
CC      ADDRESSEE: Pennie & Edmonds
CC      STREET: 1155 Avenue of the Americas
CC      CITY: New York
CC      STATE: New York
CC      COUNTRY: U.S.A.
CC      ZIP: 10036-2711
CC      COMPUTER READABLE FORM:
CC      MEDIUM TYPE: Floppy disk
CC      COMPUTER: IBM PC compatible
CC      OPERATING SYSTEM: PC-DOS/MS-DOS
CC      SOFTWARE: Patent In Release #1.0, Version #1.25
CC      CURRENT APPLICATION DATA:
CC      APPLICATION NUMBER: US/08/471,052A
CC      FILING DATE: 06-JUNE-1995
CC      CLASSIFICATION: 530
CC      ATTORNEY/AGENT INFORMATION:
CC      NAME: Mastrock, S. Leslie
CC      REGISTRATION NUMBER: 18,872
CC      REFERENCE/DOCKET NUMBER: 1101-179
CC      TELECOMMUNICATION INFORMATION:
CC      TELEPHONE: 212 790-9090
CC      TELEFAX: 212 865-8864/9741
CC      TELEX: 66141 PENNIE
CC      INFORMATION FOR SEQ ID NO: 144:
CC      SEQUENCE CHARACTERISTICS:
CC      LENGTH: 66 bases
CC      TYPE: nucleic acid
CC      STRANDEDNESS: single
CC      TOPOLOGY: unknown
CC      MOLECULE TYPE: DNA
SQ      SEQUENCE 66 BP; 2 A; 3 C; 4 G; 2 T; 55 OTHER.
Query Match              11.7%; Score 21; DB 1; Length 66;
Best Local Similarity 5.2%; Pred. No. 2,73e-01;
Matches    3; Conservative 18; Mismatches 37; Indels 0; Gaps 0;
DB        4 AGNNNNBNNBNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNC 61
QY       10 AGTGGCACCATTGGGGGTCTGGGCTGCCCTTCCTTGACCCCTTCGGCAC 67
RESULTS 11
ID     US-08-471-052A-143 STANDARD; DNA: UNC; 68 BP.
AC      xxxxxx
DE      Sequence 143, Application US/08471052A
CC      Sequence 143, Application US/08471052A
CC      Patent No. 5625033
CC      GENERAL INFORMATION:
CC      APPLICANT: Kay, B. K.
CC      APPLICANT: Fowles, D. M.
CC      TITLE OF INVENTION: Totally Synthetic Affinity Reagents
CC      NUMBER OF SEQUENCES: 166
CC      CORRESPONDENCE ADDRESS:
CC      ADDRESSEE: Pennie & Edmonds
CC      STREET: 1155 Avenue of the Americas

```

```
CC CITY: New York
CC STATE: New York
CC COUNTRY: U.S.A.
CC ZIP: 10036-2711
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patent Release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC FILING DATE: 06-JUNE-1995
CC CLASSIFICATION: 530
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-179
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: 212 790-9090
CC TELEFAX: 212 869-8864/9741
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 143:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 68 bases
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: unknown
CC MOLECULE TYPE: DNA
SQ SEQUENCE 68 BP; 3 A; 3 C; 5 G; 3 T; 54 OTHER.*

Query Match      11.7%; Score 21; DB 1; Length 68;
Best Local Similarity 13.8%; Pred. No. 2,73e-01;
Matches 9; Conservative 16; Mismatches 40; Indels 0; Gaps 0;

Db          4 CTGAANNVNNVVNNVVNNVVNNVVNNVVNNVVNNVVNNVVNNVVNNVVNNVVNNVA 63
Cp          147 CTCGAGTCATCGAGAGACTGTGCAGAAAGACCTTCACTTGCTGCAAGTCATA 88
              |||:::||||:||||:||||:||||:||||:||||:||||:||||:||||:||
Db          64 CCTGG 68
Cp          87 CCCGG 83

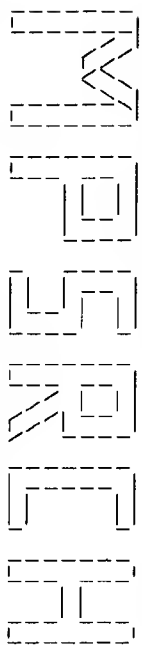
RESULT 12
ID US-08-471-052A-142 STANDARD; DNA; UNC; 69 BP.
AC xxxxxx
DT Sequence 142, Application US/08471052A
DE Sequence 142, Application US/08471052A
CC Patent No. 5625033
CC General Information:
CC APPLICANT: Kay, B. K.
CC APPLICANT: Fowles, D. M.
CC TITLE OF INVENTION: Totally Synthetic Affinity Reagents
CC NUMBER OF SEQUENCES: 166
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Pennile & Edmonds
CC STREET: 1155 Avenue of the Americas
CC CITY: New York
CC STATE: New York
CC COUNTRY: U.S.A.
CC ZIP: 10036-2711
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patentin Release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC FILING DATE: 06-JUNE-1995
CC CLASSIFICATION: 530
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistock, S. Leslie
```

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CC REGISTRATION NUMBER: 18-872
CC REFERENCE/DOCKET NUMBER: 1101-179
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: 212 790-9090
CC TELEFAX: 212 869-8864/9741
CC TELEX: 66141 PENNIE
CC INFORMATION FOR SEQ ID NO: 142:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 69 bases
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: unknown
CC MOLECULE TYPE: DNA
SQ SEQUENCE 69 BP; 2 A; 4 C; 6 G; 2 T; 55 OTHER.

Query Match      11.7%; Score 21; DB 1; Length 69;
Best Local Similarity 5.2%; Pred. No. 2,73e+01;
Matches          3; Conservative 18; Mismatches 37; Indels 0; Gaps 0

Db   7 AGNNBNNBNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNC 64
    || .....|.....|.....|.....|.....|.....|.....|
Oy   10 AAGGCCACATGGGTGCTGCCTGCCCTTGTCCTCCTTGAACCCTCGCACG 67

RESULT  13
ID     PCT-US95-11934--94 STANDARD; DNA; UNC; 74 BP.
AC     xxxxxx
DT
DI      Sequence 94, Application PC/TUS9511934
CC      Sequence 94, Application PC/TUS9511934
CC      GENERAL INFORMATION:
CC           APPLICANT: Cylogen Corporation
CC           TITLE OF INVENTION: Antigen Binding Peptides (Abltides) From
CC           TITLE OF INVENTION: Peptide Libraries
CC           NUMBER OF SEQUENCES: 103
CC           CORRESPONDENCE ADDRESS:
CC               ADDRESSER: Pennie & Edmonds
CC               STREET: 1155 Avenue of the Americas
CC               CITY: New York
CC               STATE: New York
CC               COUNTRY: USA
CC       ZIP: 10036
CC       COMPUTER READABLE FORM:
CC           MEDIUM TYPE: Floppy disk
CC           COMPUTER: IBM PC compatible
CC           OPERATING SYSTEM: PC-DOS/MS-DOS
CC           SOFTWARE: Patent In Release #1.0, Version #1.30
CC           CURRENT APPLICATION DATA:
CC               APPLICATION NUMBER: PCT/US95/11934
CC               FILING DATE: 20-SEP-1995
CC       CLASSIFICATION:
CC           ATTORNEY/AGENT INFORMATION:
CC               NAME: Mastrock, S. Leslie
CC               REGISTRATION NUMBER: 18-872
CC               REFERENCE/DOCKET NUMBER: 1101-196-228
CC           TELECOMMUNICATION INFORMATION:
CC               TELEPHONE: (212) 790-9090
CC               TELEFAX: (212) 869-9741/8864
CC               TELEX: 66141 PENNIE
CC           INFORMATION FOR SEQ ID NO: 94:
CC               SEQUENCE CHARACTERISTICS:
CC                   LENGTH: 74 base pairs
CC                   TYPE: nucleic acid
CC                   STRANDEDNESS: single
CC                   TOPOLOGY: linear
CC       MOLECULE TYPE: DNA (genomic)
CC       SEQUENCE 74 BP; 3 A; 4 C; 3 G; 1 T; 63 OTHER.
```

(TM)

Release 3.1a John F. Collins, Biocomputing Research Unit.
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MPerch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 17:26:03 1999; MasPar time 365.62 Seconds

Tabular output not generated. 1153.579 Million cell updates/sec

Title: >US-09-092-296-3
Description: (1-180) from US0902296.seq
Perfect Score: 180
N.A. Sequence: 1 CAGGAGCGCAGTCCGACCTA.....CTGAAACCTCTCCCTCT 180
Comp: GTCTCCGCTCACCGCTCAT.....GACCTTTGAGAGGAGGA

Scoring table: TABLE default
Gap 6

Mmatch STD: Dbase 0; Query 0

Searched: 2883791 segs, 1171580779 bases x 2

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database: emb1-est58
1:em_est10 2:em_est11 3:em_est17 4:em_est18 5:em_est12
6:em_est9 7:em_gss1
Database: genbank-est111
8:gb_est1 9:gb_est10 10:gb_est11 11:gb_est12 12:gb_est13
13:gb_est14 14:gb_est15 15:gb_est16 16:gb_est17
17:gb_est18 18:gb_est19 19:gb_est20 20:gb_est21
21:gb_est22 22:gb_est23 23:gb_est24 24:gb_est25
25:gb_est26 26:gb_est27 27:gb_est28 28:gb_est29
29:gb_est30 30:gb_est31 31:gb_est32 32:gb_est33 33:gb_est34
34:gb_est35 35:gb_est36 36:gb_est37 37:gb_gss1 38:gb_gss2
39:gb_gss3 40:gb_gss4 41:gb_gss5 42:gb_gss6

Statistics: Mean 9.460; Variance 1.952; scale 4.848

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description	Pred. No.
1	51	28.3	252	17	AA754459	9.68e-50
2	43	23.9	328	23	97SN1787 Rice Immature	3.33e-36
3	39	21.7	252	17	UI-R-C2P-ng-e-02-0-UI.	1.13e-29
4	31	17.2	247	17	AA754458	8.54e-16
5	30	16.7	247	17	AA754458	8.54e-16
6	28	15.6	2275	20	AF034173 Human mRNA (T	6.48e-13
7	27	15.0	2275	20	AF034173 Human mRNA (T	6.48e-13
8	23	12.8	311	11	AA325964 EST16816 Cerebellum IT	3.57e-06
9	23	12.8	459	13	AA475002 vH08h04.r1 Soares mos	3.57e-06
10	23	12.8	1287	20	AF038250 Human mRNA (T	3.57e-06

11	22	12.2	259	38	B81336	CIT-RSP-2015F16.TFC CI	6.38e-05
12	22	12.2	308	8	D40392	RIC23424 Rice shoot 0	6.38e-05
13	22	12.2	339	19	F08745	HSC1DB011 normalized	6.38e-05
14	22	12.2	427	33	N80550	za10h06.r1 Soares feta	6.38e-05
15	22	12.2	433	34	W79098	zd75h10.r1 Soares feta	6.38e-05
16	22	12.2	441	14	C28493	C28493 Rice callus cdn	6.38e-05
17	22	12.2	513	40	AQ235722	HS.2015.B2.C08.T7 CIT	6.38e-05
18	22	12.2	634	22	A1063013	CH02423 5p-time GH Dros	6.38e-05
19	22	12.2	1025	37	B12587	F22011-SP6.1 IGF Arabi	6.38e-05
20	22	11.7	243	8	D22101	RIC10342A Rice callus	1.04e-03
21	21	11.7	288	11	AA335414	EST39832 Epididymus Ho	1.04e-03
22	21	11.7	299	26	A1382671	q205106.r1 NCI_CGAP_CL	1.04e-03
23	21	11.7	299	8	M79528	WEST00065 Mxed stage,	1.04e-03
24	21	11.7	317	19	F10052	HSC396122 normalized	1.04e-03
25	21	11.7	343	8	T47417	Yb13f12.r1 Stratigene	1.04e-03
26	21	11.7	364	34	W33870	mc56b03.r1 Soares mos	1.04e-03
27	21	11.7	412	38	AQ007423	CIT-RSP-2292F15.TR CIT	1.04e-03
28	21	11.7	416	8	T08769	EST08661 Infant Brain,	1.04e-03
29	21	11.7	417	30	R36249	Yb91109.r1 Soares plac	1.04e-03
30	21	11.7	422	10	AA275959	vc27e03.r1 Barstead MP	1.04e-03
31	21	11.7	426	8	T15255	crs853 lambdaZAPST Ric	1.04e-03
32	21	11.7	437	37	FR0032574	Fugu rubripes GSS sequ	1.04e-03
33	21	11.7	451	38	AQ024376	HS.2182.AL.D09.MF.CIT	1.04e-03
34	21	11.7	468	35	AA046966	zF50a10.r1 Soares rect	1.04e-03
35	21	11.7	475	28	A1522104	t433905.r1 NCI_CGAP_KI	1.04e-03
36	21	11.7	502	15	AA586074	28723 Lambda-PR12 Arab	1.04e-03
37	21	11.7	516	31	H41536	Yp11c01.s1 Soares adu1	1.04e-03
38	21	11.7	524	10	AA239967	mw24907.r1 Soares mos	1.04e-03
39	21	11.7	533	15	AA593745	n183q11.s1 NCI_CGAP_Br	1.04e-03
40	21	11.7	557	23	A1102979	EST212268 Normalized r	1.04e-03
41	21	11.7	567	30	R61539	Yh16f01.s1 Soares inf1	1.04e-03
42	21	11.7	630	39	AO201160	RPc111-46K18.TJ RPc111	1.04e-03
43	21	11.7	694	36	AA140679	CR00462 5p-time CR Dros	1.04e-03
44	21	11.7	762	37	B19344	T2711-SP6.1 TMU Arabi	1.04e-03
45	20	11.1	492	34	W96005	ze09b02.r1 Soares_feta	1.35e-02

ALIGNMENTS

RESULT	1	AA754459	252 bp	mRNA	EST	20-JAN-1998
LOCUS		97SN1787 Rice Immature Seed	lambda ZAPII	CDNA Library	Oryza sativa	
DEFINITION		CDNA clone 97SN1787, mRNA sequence.				
ACCESSION		AA754459				
VERSION		92801165				
KEYWORDS		AA754459.1	GI:2801165			
SOURCE		EST.				
ORGANISM		Oryza sativa.				
		Oryza sativa				
		Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;				
		euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;				
		Poaceae; Oryza.				
REFERENCE		1 (bases 1 to 252)				
AUTHORS		Nahm,B.H., Kim,J.K., Cheong,J.U., Kim,S.I., Hahn,T.R., Moon,E.P.,				
		Kim,W.T., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y.,				
		Lee,M.C. and Eun,M.Y.				
TITLE		Large-scale Sequencing Analysis of ESTs from Rice Immature Seed				
JOURNAL		Unpublished (1998)				
COMMENT		On Jan 14, 1998 this sequence version replaced gi:1797457.				

Contact: Eun M.Y.
Department of Cytogenetics
National Inst. of Agri. Sci. and Tech, RDA
Suwon, Kyunggi-do, Korea
Tel: 82 331 290 0301
Fax: 82 331 290 0307
Email: myeun@sunt20.osti.re.kr
Submitted by Baek Hie Nahm, Dept of Biological Science, Myongji
University, Yongin, Korea. 449-728 bhnam@bioserver.myongji.ac.kr
Seq primer: M13 Reverse Primer.
Location/Qualifiers
1. 252
/organism="Oryza sativa"

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/cultivar="M1yang23"
/note="vector: pBluescript SK(+); Site_1: EcoRI; Site_2:
XhoI; Directional cDNA library inserted into lambda ZAPIT
vector at 5' end with EcoRI and 3' end with Xho I site."
/db_xref="taxon:4530"
/map="6"
/clone="97SN1787"
/clone.lib="Rice Immature Seed Lambda ZAPIT cDNA Library"
/tissue.type="Immature Seed"
/dev_stage="5 days after pollination"
/lab_host="E. coli SOLR"

BASE COUNT      5 a      21 c      12 g      35 t      179 others
ORIGIN

Query Match      28.3%; Score 51; DB 17; Length 252;
Best Local Similarity 12.3%; Pred. No. 9.68e-50;
Matches 20; Conservative 84; Mismatches 56; Indels 2; Gaps 2;

Db 29 BIVVCAASHGNMAYHNCIBRGTHCCKRYNMSTMTGTYMBNBSGDMHMBYBNTKVD 88
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Oy 4 GAGCGAGTGGCCACTATGAGGTCTGGCGCCCTTGTCTCTTCAACCCCTCTTGC 63
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 89 VGNHTCSMRBYTMAH-YHDYTCBRYNNNDYHMHBMRYBTGCTTMCMBHYNT 147
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Oy 64 CAGCTCAGATGACGAGCGCGGATGACTTGCAGACTGACGAGAGAGCTTTTCT 123
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 148 KCTASGHTSTNYDKS-STNTWCYBSYDKSMHCYCSBV 188
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Oy 124 GACAGTCTCTCTATGAGTCAGCTTCTGGAATCTCTGA 165
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 2
LOCUS A116523 328 bp mRNA EST 11-FEB-1999
DEFINITION UI-R-C2p-ng-e-02-0-UI.s1 UI-R-C2p Rattus norvegicus cDNA clone
ACCESSION A116523
VERSION 93637300
KEYWORDS A116523.1 GI:3637300
SOURCE EST.
ORGANISM Norway rat.
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
REFERENCE 1 (bases 1 to 328)
AUTHORS Bonaldo,M.F., Lennon,G. and Soares,M.B.
TITLE Normalization and subtraction: two approaches to facilitate gene
discovery
JOURNAL Genome Res. 6 (9), 791-806 (1996)
MEDLINE 97044477
COMMENT On Jan 14, 1998 this sequence version replaced gi:1877567.

Contact: Soares, MB
Program for Rat Gene Discovery and Mapping
University of Iowa
451 Einstein Medical Research Building Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: msoares@blue.weeg.uiowa.edu
The sequence tag present in the cDNA between the NotI site and the
oligo-dT track served to identify it as a clone from the normalized
adult lung library. cDNA Library Preparation: M. Fatima Bonaldo,
Ph.D. Clone distribution: clones will be available through Research
Genetics
Seq primer: M13 forward.
Location/Qualifiers
1..328
/organism="Rattus norvegicus"
/strain="Sprague-Dawley"
/note="vector: p773p-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; The UI-R-C2p
library is a subtracted library derived from the UI-R-C1
library, which is a subtracted library derived from the
UI-R-C0 library. The UI-R-C0 library consisted of a

```

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mixture of individually tagged normalized libraries
constructed from rat placenta, adult lung, brain, liver,
kidney, heart, spleen, ovary, muscle, 8, 12 and 18-day
embryo. The tag is a string of 3-5 nucleotides present
between the Not I site and the oligo-dT track which allows
identification of the library of origin of a clone within
the mixture. The subtracted library (UI-R-C2p) was
constructed as follows: PCR amplified cDNA inserts from
UI-R-C1 clones from which 3' ESTs had been derived was
used as a driver in a hybridization with the UI-R-C1
library. In the form of single stranded circles. The
remaining single-stranded circles (subtracted library) was
purified by hydroxyapatite column chromatography,
converted to double-stranded circles and electroporated
into DH10B bacteria (Life Technologies) to generate the
UI-R-C2p library. This procedure has been previously
described (Bonaldo, Lennon and Soares, Genome Research 6:
791-806, 1996)"
/db_xref="taxon:10116"
/clone="UI-R-C2p-ng-e-02-0-UI"
/clone.lib="UI-R-C2p"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies)"

BASE COUNT      62 a      77 c      98 g      91 t
ORIGIN

Query Match      23.9%; Score 43; DB 23; Length 328;
Best Local Similarity 73.6%; Pred. No. 3.33e-36;
Matches 67; Conservative 0; Mismatches 24; Indels 0; Gaps 0;

Db 210 AGAGCGAGATCTTTGGAGCATGTCGAGAACCGGAGCTTGGAGCTTGGCCCTGA 269
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 180 AGAGCGAGATCTTTGACGATTCGACGAGAGCTGACGAGAGCTTGGAGCA 121
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 270 GATCGCTCGTCACTTCTTCACTTCAAGAGTCA 300
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Cp 120 AAGAGCTCTTCACTTCACTTCAAGAGTCA 90
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 3
LOCUS AA754459 252 bp mRNA EST 20-JAN-1998
DEFINITION 97SN1787 Rice Immature Seed Lambda ZAPIT cDNA library Oryza sativa
ACCESSION AA754459
VERSION 92801165
KEYWORDS AA754459.1 GI:2801165
SOURCE EST.
ORGANISM Oryza sativa.
Oryza sativa
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
Poaceae; Oryza.
REFERENCE 1 (bases 1 to 252)
AUTHORS Nahm,B.H., Kim,J.K., Cheong,J.Y., Kim,S.I., Hahn,T.R., Moon,E.P.,
Kim,W.T., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y.,
Lee,M.C. and Eun,M.Y.
TITLE Large-scale Sequencing Analysis of ESTs from Rice Immature Seed
unpublished (1998)
COMMENT On Jan 14, 1998 this sequence version replaced gi:197457.

Contact: Eun M.Y.
Department of Cyto genetics
National Inst. of Agri. Sci. and Tech, RDA
Suwon, Kyunggi-do, Korea
Tel: 82 331 290 0301
Fax: 82 331 290 0307
Email: myeun@snu20.asi.re.kr
Submitted by Baek Hie Nahm, Dept of Biological Science, Myongji
University, Yongin, Korea, 449-728 binahm@josever.myongji.ac.kr
Seq primer: M13 Reverse Primer.
Location/Qualifiers
1..252
/organism="Oryza sativa"

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FEATURES
source
1..252
/organism="Oryza sativa"

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ORIGIN	Query Match	Best Local	Matches	Score	DB	Length
17.2%	Score 31;	DB 17;	Length 247;			
8.3%	Pred. No. 2.88e-17;					
11;	Conservative	71;	Mismatches 49;	Indels 2;	Gaps 2;	
100	VNHSQANNRCSNVVYVWELTACVBYHDBRAHYDTRCINDGKONTASDNGTSAT	159				
174	CACAGCTTTTACACATTTCCAGAGAACTGACATCATTAGAGAACTGTCAGAAAGC	115				
160	KRTGCTDITDSCGGGCKRKYVYSSBYRCGVNMYFTTSMATDKTBSM-DMSRRS	218				
114	TCCTTACGCTTCACTTGCAGAAAGT-CATACCGGCGCTTCATGTAGCTGCAAGAG	56				
219	RHYGRNMBNKRK	231				
55	GGTCAAGAGAGG	43				
RESULT	5	AA754458	247 bp	mRNA	EST	20-JAN-1998
LOCUS	97SN1784	Rice	Immature Seed	Lambda ZAPII	CDNA Library	Oryza sativa
DEFINITION	CDNA clone 97SN1784, mRNA sequence.					
ACCESSION	AA754458					
NID	92801164					
VERSION	AA754458.1	GI:2801164				
KEYWORDS	EST.					
SOURCE	Oryza sativa.					
ORGANISM	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Oryza.					
REFERENCE	1 (bases 1 to 247)					
AUTHORS	Nahm,B.H., Kim,J.K., Cheong,J.J., Kim,S.I., Hahn,T.R., Moon,E.P., Kim,W.T., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y., Lee,M.C. and Eun,M.Y.					
TITLE	Large-scale Sequencing Analysis of ESTs from Rice Immature Seed					
JOURNAL	Unpublished (1998)					
COMMENT	On Jan 14, 1998 this sequence version replaced gi:1797455.					
FEATURES	source	1..247				
	/organism="Oryza sativa"					
	/cultivar="Milyang23"					
	/note="Vector: pBluescript SK(+); site_1: EcoRI; site_2: XhoI; Directional cDNA library inserted into lambda ZAPII vector at 5' end with EcoRI and 3' end with Xho I site."					
	/db_xref="taxon:4530"					
	/map="6"					
	/clone="97SN1784"					
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	/dev_stage="5 days after pollination"					
	/lab_host="E. coli SOLR"					
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	/map="6"					
	/clone="97SN1784"					
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	/map="6"					
	/clone="97SN1784"					
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	/dev_stage="5 days after pollination"					
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	/map="6"					
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	/tissue_type="Immature Seed"					
	/dev_stage="5 days after pollination"					
	/lab_host="E. coli SOLR"					
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y 32 CCGCCCCCTTGACCTCTCTTGGACCGCTCTTGACGAGTCACATGGAACAGGCGCGGATATG 91
Db 171 DCGGGCMKRYATVSSSBRCGVNMMVTSMJTDKSTKMSBMSRRSRVHYGFRMBKK 230
y 92 ACTTTGGCAGCAGGATGAGAGATCTTTTCTGACAAAGTCTCTTATGATGTCAGCTTC 151
Db 231 RGMSRNMTDRTK 242
y 152 CTGGAATTGCTT 163

RESULT 6
LOCUS AF034173 2275 bp mRNA EST 30-MAR-1998
DEFINITION AF034173 Human mRNA (Tripodis and Ragousis) Homo sapiens cDNA
ACCESSION AF034173
NID 92707735
KEYWORDS AF034173.1 GI:2707735
SOURCE EST.
ORGANISM human.
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 2275)
AUTHORS Tripodis,N and Ragousis,J.
TITLE Generation of a transcription map in the region immediately
centromeric to human MHC across the 6p21.2-6p21.3 chromosomal
boundary
JOURNAL Unpublished (1997)
COMMENT On Jan 19, 1998 this sequence version replaced gi:2045115.

JOURNAL
COMMENT

FEATURES
SOURCE location/Qualifiers
1..2275
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="6p21.3"
/clone="ntcon2 contig"
/clone_1ib="Human mRNA (Tripodis and Ragousis)"
BASE COUNT 438 a 619 c 470 g 599 t 149 others
ORIGIN
Query Match 15.64; Score 28; DB 20; Length 2275;
Best Local Similarity 12.34; Pzed. No. 6.48e-13;
Matches 9; Conservative 44; Mismatches 19; Indels 1; Gaps 1
Db 1523 RCGKCKCMKRYKRYKRYSTYKSKMSRWYTTTWTGCKCT-SMKASACAMRMKMGKGS 1501
Cp 145 GGCACCTCAGAGGAGGAGACTTTCAGAAAGACTCTCTACCTGACATGTCAAAGTCAATACC 86
Db 1582 RSSRSRYGMYGSM 1594
Cp 85 CGGCCCTGTTC 73

RESULT 7
LOCUS AF034173 2275 bp mRNA EST 30-MAR-1998
DEFINITION AF034173 Human mRNA (Tripodis and Ragousis) Homo sapiens cDNA
ACCESSION AF034173
NID 92707735
KEYWORDS AF034173.1 GI:2707735
SOURCE EST.
ORGANISM human.
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 2275)
AUTHORS Tripodis,N and Ragousis,J.

```

TITLE
Generation of a transcription map in the region immediately centromeric to human MHC across the 6p21.2-6p21.3 chromosomal boundary
Unpublished (1997)
On Jan 19, 1998 this sequence version replaced gi:2045115.

JOURNAL COMMENT
Contact: Tripodis, Nikos
Division of Medical and Molecular Genetics
Guys Hospital
7th floor, Guy's tower, London SE1 9RT, UK
Email: nikos@nki.ni.
Location/Qualifiers
1..2275
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="6p21.3"
/clone="ntcon2 contig"
/clone_1lb="Human mRNA (Tripodis and Ragoussis)"
438 a 619 c 470 g 599 t 149 others

BASE COUNT
438 a 619 c 470 g 599 t 149 others

ORIGIN
Query Match 15.0%; Score 27; DB 20; Length 2275;
Best Local Similarity 17.8%; Pied. No. 1,64e-11;
Matches 19; Conservative 53; Mismatches 33; Indels 2; Gaps 2;
Db 1534 YKRYSTYYSKMSRWYTYTYTYTTCCTCMKASACAMRMKMGSSRSRSGYWGCS 1593
OY 39 TTGTCCTCCCTGACACCTCCTGAGCAGCTCAGTACATGACAGGCGGATGACTTTC 98
OY 99 AA-CTGAAGCTGAAGAGACTCTTTCCTGA-CAAGTTCCTCATAGAT 143

RESULT 8
AA323964 311 bp mRNA EST 20-APR-1997
LOCUS EST26816 Cerebellum II Homo sapiens cDNA 5' end, mRNA sequence.
ACCESSION AA323964
NID 91976290
VERSION AA323964.1 GI:1976290
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (bases 1 to 311)
Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A.,
Bult,C.D., Lee,N.H., Kitzness,E.F., Weinstock,K.G., Gocayne,J.D.,
White,O., Sutton,G., Blake,J.A., Brannon,R.C., Maniatis,C.,
Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,
Fitzgerald,L.M., Fitzhugh,W.M., Fritchman,J.L., Geoghagen,N.S.,
Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S., Jr.,
Kelleys,J.M., Kelley,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M.,
Moreno-Valenzuela,R.F., McDonald,L.A., Naucek,D.T., Pelligrino,S.M.,
Phillips,C.A., Ryder,S.E., Scott,T.R., Saudek,D.M., Shiley,R.,
Smell,K.V., Spriggs,T.A., Uterback,T.R., Weidman,J.F., Li,Y.,
Bendall,K.D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,
Dirke,D., Feng,D.-F., Ferrrie,A., Fischer,C., Hastings,G.A., K.,
He,W.W., Hu,J.S., Greene,J.W., Gruber,J., Hudson,P., Kim,A.K.,
Kozak,D.L., Kunsch,C., Hungjun,J., Li,H., Meisner,P.S., Olsen,H.,
Raymond,L., Wei,Y.F., Wang,J., Xu,C., Yu,G.L., Ruben,S.M.,
Dillion,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C.,
Fraser,C.M. and Venter,J.C.
Initial assessment of human gene diversity and expression patterns
based upon 83 million nucleotides of cDNA sequence
Nature 377 (6547 suppl), 3-174 (1995)
On Apr 14, 1993 this sequence version replaced gi:693635.

TITLE
JOURNAL
MEDLINE
COMMENT
Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA

RESULT 11
 LOCUS B81136 259 bp DNA GSS 24-JUN-1998
 DEFINITION CIT-HSP-2015F16.TFC CIT-HSP Homo sapiens genomic clone 2015F16,
 genomic survey sequence.
 ACCESSION B81136
 NID 92868159
 VERSION B81136.1 GI:2868159
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
 Primates; Catarrhini; Hominoidea; Homo.
 REFERENCE 1 (bases 1 to 259)
 Adams,M.D., Rounsley,S.D., Field,C.E., Bass,S., Linher,K.,
 Golden,K., Berry,K., Granger,D., Suh,E., White,C., Shizuya,H.,
 Simon,M. and Venter,J.C. Genenger,D., Suh,E., White,C., Shizuya,H.,
 Use of a random BAC End Sequence Database for Sequence-Ready Map
 Building
 TITLE Use of a random BAC End Sequence Database for Sequence-Ready Map
 Building
 JOURNAL Unpublished (1997)
 COMMENT
 Contact: Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: mdadams@tigr.org
 Clones are available from Research Genetics (info@resgen.com). BAC
 end search page:
 http://www.tigr.org/db/humgen/Bac_end_search/Bac_end_search.html
 Seq primer: M13-21
 Class: BAC ends.
 FEATURES
 source location/Qualifiers
 1..259
 /organism="Homo sapiens"
 /note="Vector: pheb10AC11, site_1: HindIII, site_2:
 HindIII"
 /db_xref="taxon:9606"
 /clone="2015F16"
 /clone_1b="CIT-HSP"
 /sex="Male"
 /cell_type="Sperm"
 BASE COUNT 68 a 43 c 56 g 92 t
 ORIGIN
 Query Match 12.2%; Score 22; DB 38; Length 259;
 Best Local Similarity 89.3%; Pred. No. 6.38e-05;
 Matches 25; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 DB 59 GCTTCTGGATTCTTGAATCGCTG 86
 OY 147 GCTTCTGGATTCTTGAATCGCTG 174
 RESULT 12
 LOCUS D40392 308 bp mRNA EST 11-NOV-1994
 DEFINITION R16S2342A Rice shoot Oryza sativa cDNA, mRNA sequence.
 ACCESSION D40392
 NID 6569543
 VERSION D40392.1 GI:569543
 KEYWORDS EST.
 SOURCE Oryza sativa.
 ORGANISM Oryza sativa
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
 Poaceae; Oryza.
 REFERENCE 1 (bases 1 to 308)
 Sasaki,T., Miyao,A. and Yamamoto,K.
 Rice cDNA from callus 1995
 TITLE Rice cDNA from callus 1995
 JOURNAL Unpublished (1995)
 COMMENT
 Contact: Takuji Sasaki
 National Institute of Agrobiological Resources

Rice Genome Research Program
 2-1-2 Kannondai, Tsukuba
 Ibaraki,
 Japan 305
 Tel: 0298-38-7441
 Fax: 0298-38-7468
 Email: tsasaki@abr.affrc.go.jp.
 FEATURES
 source location/Qualifiers
 1..308
 /organism="Oryza sativa"
 /strain="Nipponbare, sub-species japonica"
 /note="Etolated shoot (8 days old)"
 /db_xref="taxon:4530"
 /clone_1b="Rice shoot"
 BASE COUNT 57 a 126 c 57 g 68 t
 ORIGIN
 Query Match 12.2%; Score 22; DB 8; Length 308;
 Best Local Similarity 73.9%; Pred. No. 6.38e-05;
 Matches 34; Conservative 0; Mismatches 12; Indels 0; Gaps 0;
 DB 115 GCTGCACCTCTCTCTCTCTCTCTCTCAAGAGCTCCATGGA 160
 OY 31 GCTGCCCTTGCTCTCTCTGACCTCTTGCGACGCTCAGCATGGA 76
 RESULT 13
 LOCUS F08745 339 bp mRNA EST 20-FEB-1995
 DEFINITION HSC1DB011 normalized infant brain cDNA Homo sapiens cDNA clone
 c-1db01, mRNA sequence.
 ACCESSION F08745
 NID 9672165
 VERSION F08745.1 GI:672165
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominoidea; Homo.
 REFERENCE 1 (bases 1 to 339)
 Auffray,C., Behar,G., Bois,F., Bouchier,C., da Silva,C.,
 Devignes,M.D., Duprat,S., Houlgatte,R., Jumeau,M.N., Lamy,B.,
 Lorenzo,F., Mitchell,H., Mariage-Samson,R., Pletu,G., Pouillot,Y.,
 Sebastiani-Kabatchis,C. and Tessier,A.
 IMAGE: molecular integration of the analysis of the human genome
 and its expression
 TITLE and its expression
 JOURNAL C. R. Acad. Sci. III, Sci. Vie 318 (2), 263-272 (1995)
 MEDLINE 95277534
 COMMENT On Sep 21, 1992 this sequence version replaced gi:276248.
 Contact: Genethon
 Genexress-Genethon
 Genethon Centre de recherche sur le Genome Humain
 1,rue de l'Internationale, BP60 91002 EVRY Cedex, FRANCE
 Tel: 33169472800
 Fax: 33160778698
 Email: genexress@genethon.fr
 Single read.
 Genexpress_library_id: G; Genexpress_sequence_id: y3c-1db01
 Seg primer: (-21)M13-universal
 High quality sequence stop: 277.
 FEATURES
 source location/Qualifiers
 1..339
 /organism="Homo sapiens"
 /note="Organ: Brain; Vector: lambda B; Site_1: HindIII;
 site_2: NotI; sex=Female; dev_stage=3 months old;
 isolate=muscular atrophy patient; tissue_type=total
 brain; total mRNA was oligo-(dT) primed and directionally
 cloned 5' -> 3' into the HindIII -> NotI sites of the
 lambda B vector. Clone library from B.Souares, Psychiatry
 Dept. Columbia University, USA. Normalization_method:
 Bento Soares, P.N.A.S in press"
 /db_xref="taxon:9606"
 /clone="c-1db01"

Mon Oct 25 11:54:02 1999

US-09-092-296-3.rst

Page 8

36 CCCTGTCTCTCTTGACCTCTCTGGCAGCTCACATGG 75

Search completed: Sun Oct 24 17:32:17 1999
Job time : 374 secs.


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OY 1 MGGSLPVLVLLTLGGSHGPEMTLQLEKESFLTNSSTESFLELLKLLHLPSG 60
DB 61 tsytlharsqghvvcnt 78
OY 61 TSYTLHARSQGHVVCNT 78

RESULT 2
ID W0327 standard; Protein: 46 AA.
AC W0327;
DE 21-DEC-1998 (first entry)
DE Secreted protein FB78_1.
KW Secreted protein; D0123_1; human.
OS Homo sapiens.
FH Key Location/Qualifiers
FT Peptide 8..20
FT /note= "predicted leader/signal sequence, or
FT transmembrane domain"
FN W09838209-A2.
PD 03-SEP-1998.
PE 25-FEB-1998; U03697.
PR 24-FEB-1998; US-028724.
PR 26-FEB-1997; US-805819.
PA (GEMT ) GENETICS INST INC.
PI Acostino MJ, Jacobs K, Lavallie ER, McCoy JM, Werberg D,
PI Racine LA, Spaulding V, Treacy M;
PI WPI: 98-481138/41.
DR N-PSDB: V33199.
DE New isolated polynucleotide(s) and encoded polypeptide(s) -
DE obtained from human foetal kidney, adult colon, adult brain,
DE brain and placenta cDNA libraries.
PS Claim 36; Page 83; 103pp; English.
CC This is the amino acid sequence of novel human secreted protein
CC FB78_1, as deduced from a full-length cDNA clone (see V33199)
CC obtained from a human adult placenta cDNA library. Database
CC searching revealed some similarity between FB78_1 and some known
CC sequences. The invention provides new isolated polynucleotides
CC (see V33199-99), from human foetal kidney, adult colon, adult brain,
CC foetal brain and placenta cDNA libraries, that code for secreted
CC proteins (see W0319-27). The clones can be used for recombinant
CC production of the polypeptides, which may have activities such as
CC e.g. nutritional activity, cytokine and cell proliferation or
CC differentiation activity, immunostimulant or immunosuppressive,
CC haematopoiesis regulating activity, tissue growth activity, activin
CC or inhibin activity, chemotactic or chemokinetic activity,
CC haemostatic and thrombolytic activity, receptor/ligand activity,
CC antiinflammatory activity, cadherin/tumour invasion suppressor
CC activity, tumour inhibition activity, or other activities.
SQ Sequence 46 AA.

Query Match 15.4%; Score 86; DB 36; Length 46;
Best Local Similarity 56.5%; Pred. No. 3.89e+01;
Matches 13; Conservative 6; Mismatches 3; Indels 1; Gaps 1;

DB 6 gaalptllllalrctfngargp 28
OY 2 GSGCLP-LVLLTLGLSSHGTCGPG 23

RESULT 3
ID W73408 standard; Protein: 47 AA.
AC W73408;
DE 19-FEB-1999 (first entry)
DE Human secreted protein encoded by Gene No. 12.
DE Secreted protein; human; protein therapy; gene therapy; blood disorder;
DE pathological condition; diagnosis; cancer; neurological disorder;
DE developmental abnormality; foetal deficiency; leukemia; hepatic disease;
DE immune system disorder; Alzheimer's disease; cognitive disorder;
DE schizophrenia; prostate disease; autoimmune disorder; AIDS.
OS Homo sapiens.
FH Key Location/Qualifiers
FT MISC.difference 47

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FN W09854206-A1.
PD 03-DEC-1998.
PE 28-MAY-1998; U10868.
PR 29-AUG-1997; US-056296.
PR 30-MAY-1997; US-044039.
PR 30-MAY-1997; US-048093.
PR 30-MAY-1997; US-048101.
PR 30-MAY-1997; US-048190.
PR 30-MAY-1997; US-048356.
PR 30-MAY-1997; US-050925.
PR 29-AUG-1997; US-056250.
PR 29-AUG-1997; US-056293.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Carter KC, Dillon PJ, Endress GA, Feng P, Ni J,
PI Rosen CA, Ruben SM, Yu G;
PI WPI: 99-070209/06.
DR N-PSDB: V08832.
DE New isolated human genes - useful for diagnosis and treatment of,
DE e.g. cancers, neurological disorders, immune diseases, developmental
DE disorders or blood disorders
DE Claim 11; Page 152-153; 188pp; English.
PS This sequence is encoded by a cDNA of the invention, designated
CC Gene No. 12. This sequence represents a human secreted protein, and is
CC expressed in activated neutrophils, endothelial cells, T-cells and
CC to a lesser extent in brain and liver.
CC The DNA sequences of the invention and their corresponding secreted
CC polypeptides are useful for prevention, treating or ameliorating medical
CC conditions, e.g. by protein or gene therapy. Also pathological conditions
CC can be diagnosed by determining the amount of the new polypeptides in a
CC sample or by determining the presence of mutations in the DNA sequences.
CC Specific uses are described for each of the DNA sequences and the encoded
CC proteins, based on which tissues they are most highly expressed in, and
CC include developing products for the diagnosis or treatment of cancer,
CC tumors, neurological disorders, developmental abnormalities and foetal
CC deficiencies, blood disorders, leukemias, diseases of the immune system
CC (including allergies or asthma), hepatic disease, Alzheimer's and
CC cognitive disorders, schizophrenia, prostate diseases, autoimmune
CC disorders and AIDS. The polypeptides are also useful for identifying
CC their binding partners.
SQ Sequence 47 AA;

Query Match 15.4%; Score 86; DB 38; Length 47;
Best Local Similarity 56.5%; Pred. No. 3.89e+01;
Matches 13; Conservative 6; Mismatches 3; Indels 1; Gaps 1;

DB 6 gaalptllllalrctfngargp 28
OY 2 GSGCLP-LVLLTLGLSSHGTCGPG 23

RESULT 4
ID W02702 standard; peptide: 325 AA.
AC W02702;
DE 13-NOV-1996 (first entry)
DE G-protein coupled bovine adrenal angiotensin II type-1 receptor.
DE G-protein coupled receptor; ligand binding assay; transmembrane domain;
DE schizophrenia; dopamine; cAMP; adenosine; throbmin; adenylytic; opsin;
DE muscarinic acetylcholine; endothelin; bombesin; endocrine; rhodopsin;
DE odorant; cytomagalovirus; serotonergic.
OS Bos taurus.
FH Key Location/Qualifiers
FT 16-APR-1996.
PD 16-APR-1996.
PE 10-SEP-1992; 943236.
PR 10-SEP-1992; US-943236.
PR 09-SEP-1993; US-118270.
PA (UYNV ) UNIV NEW YORK STATE.
PI Murphy RB, Schuster DI;
PI WPI: 96-208785/21.
DE New dopamine receptor peptide - useful as antipsychotic agent, e.g.
DE for treating schizophrenia
DE Dislosure; Column 128-132; 184pp; English.
PS Proteins W02657-W02730 represent a range of G-protein coupled receptor

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CC (GPR) proteins selected from cAMP, adenosine, muscarinic acetylcholine, CC adrenergic, thrombin, endothelin, bombesin, endocrine, rhodopsin, opsin, CC odorant, cytomagaloviral and other GPR proteins. The receptor proteins CC were used to design polypeptides, pref. based on the transmembrane CC domains, for use in G-protein coupled receptor ligand binding assays. CC The polypeptide fragments retain biological activity such as binding a CC GPR ligand or modulating GPR ligand binding to a GPR (see W02747-W02999 CC for examples of polypeptide fragments). The polypeptide fragments can CC be used in compositions for treating subjects suffering from a pathology CC related to a GPR abnormality e.g. a psychotic disorder such as CC schizophrenia.

CC Sequence 325 AA:

Query Match 15.2%; Score 85; DB 19; Length 325;
Best Local Similarity 44.4%; Pred. No. 4.56e+01;
Matches 12; Conservative 5; Mismatches 8; Indels 2; Gaps 2;

Db 27 ymkltkyasvflnaladictfltlp 53
34 FLNNSYESSFL-EL-LEKICLULHLP 58

RESULT 5
ID R48730 standard; Protein; 325 AA.
AC R48730;
DE 06-JUN-1996 (first entry)
DT G-protein coupled bovine adrenal angiotensin II type-1 receptor protein.
KW G-protein coupled receptor; ligand binding assay; transmembrane domain;
KW psychotic disorder; schizophrenia; dopamine; cAMP; adenosine; thrombin;
KW muscarinic acetylcholine; adrenergic; endothelin; bombesin; endocrine;
KW rhodopsin; opsin; odorant; cytomagalovirus.
OS Bos taurus.
PN W09405695-A1.
PD 17-MAR-1994.
PR 09-SEP-1993; U08528.
PR 10-SEP-1993; US-943236.
PA (UNIV) UNIV NEW YORK STATE.
PI Murphy RB, Schuster DI;
DR WPI; 94-101120/12.
PT Polypeptides of G-coupled receptor proteins (GPRs) - useful for
PT binding GPR ligands or modulating GPR binding
PS Disclosure; Page 106-107; 160pp; English.
CC Proteins R48685-R48758 represent a range of G-protein coupled receptor
CC proteins selected from cAMP, adenosine, muscarinic acetylcholine,
CC adrenergic, thrombin, endothelin, bombesin, endocrine, rhodopsin, opsin,
CC odorant, cytomagaloviral and other G-protein coupled receptors. The
CC receptor proteins were used to design polypeptides, pref. based on the
CC transmembrane domains, for use in G-protein coupled receptor ligand
CC binding assays. The polypeptide fragments retain biological activity
CC such as binding a GPR ligand or modulating GPR ligand binding to a GPR
CC (see R48759-R48758, R50569-R50807 and R89189-R89195 for examples of
CC polypeptide fragments). The polypeptide fragments can be used in
CC compositions for treating subjects suffering from a pathology related to
CC a GPR abnormality e.g. a psychotic disorder such as schizophrenia.
CC Sequence 325 AA:

Query Match 15.2%; Score 85; DB 16; Length 325;
Best Local Similarity 44.4%; Pred. No. 4.56e+01;
Matches 12; Conservative 5; Mismatches 8; Indels 2; Gaps 2;

Db 27 ymkltkyasvflnaladictfltlp 53
34 FLNNSYESSFL-EL-LEKICLULHLP 58

RESULT 6
ID R47338 standard; Protein; 194 AA.
AC R47338;
DE 01-JUL-1994 (first entry)
DE Peptide fragment of tetracycline transporter protein.
KW Vesicle membrane transport protein; gene therapy; screening;
KW Parkinsons disease; neurotoxin; identification; detection;
KW antibody; probe; chromaffin granule amine transporter protein;

KW resistance; tetracycline.
OS Escherichia coli.
PN W09325699-A.
PD 23-DEC-1993.
PR 11-JUN-1993; U05704.
PR 11-JUN-1992; US-899074.
PR 30-JUL-1992; US-923096.
PA (REGC) UNIV CALIFORNIA.
PI Edwards RH;
DR WPI; 94-007556/01.
PT New mammalian vesicle membrane transport protein - and corresp.
PT DNA, vectors, transformed cells and antibodies, for diagnosis and
PT treatment of neurological disorders, e.g. Parkinson disease
PS Example 2; Page 111-112; 181pp; English.
CC This sequence of the tetracycline transporter protein showed a
CC definite homology with the chromaffin granule amine transport
CC protein (CGAT) of rat. The cDNA encoding the chromaffin granule
CC amine transport protein is useful in gene therapy and as a probe
CC for detecting genomic sequences. The transport protein is used for
CC screening cytotoxic compounds implicated in Parkinsons disease,
CC diseases associated with activity of neurotoxins or psychiatric
CC disorders and to identify compounds which selectively inhibit or
CC activate its action. Antibodies raised against the transport
CC protein are useful as immunoassay reagents for detecting the protein
CC and as affinity reagents for purification.
CC Sequence 194 AA:

Query Match 14.9%; Score 83; DB 9; Length 194;
Best Local Similarity 31.0%; Pred. No. 6.23e+01;
Matches 18; Conservative 13; Mismatches 23; Indels 4; Gaps 4;

Db 1 mmsckialvltllldam-giglmvptllre-flasedlanhfyvla-lyalnvy 55
1 MGSGLPLVLLTLGSHRGCGMT-LQALKTESFLNNSYESSFLLEKICLULHLP 57

RESULT 7
ID W57046 standard; Protein; 411 AA.
AC W57046;
DE 20-AUG-1998 (first entry)
DE Mouse apoptosis inducing receptor.
KW Apoptosis inducing receptor; AIR protein; mouse; cell death regulator;
KW Type I transmembrane protein; tumour cell death; autoimmune disease;
KW therapy.
OS Mus sp.
PN W09814565-A1.
PD 05-APR-1998.
PR 03-OCT-1997; U17876.
PR 04-OCT-1996; US-044456.
PA (IMMW) IMMUNEX CORP.
PI Perkins PA;
DR WPI; 98-240077/21.
DR N-PSDB; V28701.
PT DNA encoding apoptosis inducing receptor - which is Type I
PT transmembrane protein, useful for regulating cell death
PS Claim 16; Page 33-35; 45pp; English.
CC This sequence is the mouse apoptosis inducing receptor (AIR) of the
CC invention. AIR is a Type I transmembrane protein, soluble forms of which
CC can be used to regulate cell death in a therapeutic setting. Soluble AIR
CC can also be used in vitro to block apoptosis or AIR-expressing cells, or
CC to screen agonists or antagonists of AIR activity. The cytoplasmic domain
CC of AIR can be used to develop assays for inhibitors of AIR-induced cell
CC death, which is useful to regulate cell death in a therapeutic setting as
CC well as in vitro. Agonists of AIR activity can be used to kill tumour
CC cells that express AIR, or T cells expressing AIR in autoimmune diseases.
CC Sequence 411 AA:

Query Match 14.9%; Score 83; DB 31; Length 411;
Best Local Similarity 57.1%; Pred. No. 6.23e+01;
Matches 12; Conservative 5; Mismatches 3; Indels 1; Gaps 1;

Db 16 lplllllllllllyggqggg-gms 35
lll:lll ll :lll ll :

OY 5 LPLVLLTLGSSHGTCPCMT 25

RESULT 8
ID R26954 standard; Protein: 110 AA.
AC R26954;
DT 10-FEB-1993 (first entry)
DE Human T lymphocyte receptor V-alpha22 segment.
KW TCR; IGR a 12; variable region; Immune system modulation;
OS Homo sapiens.
PN MO9213949-A.
PD 20-AUG-1992.
PF 07-FEB-1992; F00111.
PR 08-FEB-1991; FR-001487.
PR 12-APR-1991; FR-004527.
PA (ROUS) ROUSSEL-UCIAF.
PI Ferradini L, Herceud T, Roman-Roman S, Triebel F;
DR WPI: 92-300035/36.
DR N-PSDB: Q28129.
PT Nucleotide sequences and their monoclonal antibodies and
oligo-nucleotide primers - encode variable alpha-chain regions of
human T-lymphocyte receptors, for studying immune responses and
for therapy
PS Claim 8: Page 35; 65pp; French.
CC RNA was isolated from peripheral lymphocytes and converted to cDNA.
CC The cDNA was amplified by anchored PCR using C-alpha and polyC
primers, then amplified again using a different C-alpha specific
CC primer. The amplified product was SacII-restricted, inserted into
CC Bluescript SK+ vector and used to transform E.coli XL7-blue.
CC Transforms were screened with a C-alpha specific probe and DNA
CC from positive clones was sequenced in the C-alpha region. The
CC sequence designated "IGR a 12" comprises the complete coding region
CC of a gene of the subfamily V alpha 22; this subfamily was
CC previously identified by the partial sequence (113bp) AC9 (Klein M.H.,
CC et al., Proc.Natl.Acad.Sci. USA 84:6884, 1987). The peptide encoded
CC by it can be used to block specific T cell epitopes or in vaccines.
CC See Q28120-Q28172.
SQ Sequence 110 AA;
Query Match 14.7%; Score 82; DB 5; Length 110;
Best Local Similarity 34.0%; Pred. No. 7.28e+01;
Matches 18; Conservative 12; Mismatches 18; Indels 5; Gaps 5;
Db 1 myspgylvslllllgrtgdsvtqmgvrlseeflctactatgypelf 53
OY I MGSGLPLV-LLLTLGSSHGTCG-PGMTLQLKKE-SFLT-NSSY-ESSFELL 48
RESULT 9
ID W21674 standard; Protein: 169 AA.
AC W21674;
DT 29-SEP-1997 (first entry)
DE Human mitochondrial electron transport chain subunit CIT-3.
KW Mammalian artificial chromosome; MAC; selectable marker; CIT-3;
KW mitochondrial electron transport chain complex II.
OS Homo sapiens.
PN MO9716533-A1.
PD 09-MAY-1997.
PR 28-OCT-1996; U17476.
PR 31-OCT-1995; US-550717.
PA (REBC) UNIV CALIFORNIA.
PI Scheffler IE;
DR WPI: 97-272103/24.
DR N-PSDB: T72466.
PT New mammalian artificial chromosomes - comprising a mammalian
PT centromere and a unique cloning site, used for stable expression of
PT large fragments of DNA
PS Disclosure; Page 54-55; 71pp; English.
CC CIT-3 (W21674) comprises a subunit of complex II of the human
CC mitochondrial electron transport chain. It is the expression
CC product of the CIT-3 gene identified in mammalian artificial
CC chromosome MAC-8.2.3, which is contained in the human-hamster

CC hybrid cell line XEM8.2.3 (ATCC CRL 11991). A portion of the
CC CIT-3 gene (see also T72461-65) or of CIT-3 cDNA (T72466) can be
CC utilized as a unique cloning site and selectable marker in an MAC,
CC allowing the site-specific integration of an exogenous nucleic acid
CC sequence into the MAC. The MAC can be used for stable expression
CC of large fragments of DNA and also for the production of transgenic
CC animals.
SQ Sequence 169 AA;

Query Match 14.7%; Score 82; DB 23; Length 169;
Best Local Similarity 28.0%; Pred. No. 7.28e+01;
Matches 14; Conservative 20; Mismatches 13; Indels 3; Gaps 3;

Db 62 alpamsichrgtalaagsyl-fgms-allpafes-yelavslcl 108
OY 4 GLPVLTLGSSHGTCPCMTQLKKEFLNLSYSSSTLEKLCL 53

RESULT 10
ID W59924 standard; Protein: 401 AA.
AC W59924;
DT 07-DEC-1998 (first entry)
DE Human 7-transmembrane receptor HNFY20.
KW HNFY20: G-protein coupled receptor; human; infection; HIV; pain;
KW cancer; anorexia; asthma; Parkinson's disease; acute heart failure;
KW hypertension; hypertension; urinary retention; osteoporosis;
KW angina pectoris; myocardial infarction; ulcer; allergy;
KW benign prostatic hypertrophy; psychosis; anxiety; schizophrenia;
KW manic depression; delirium; dementia; mental retardation;
KW dyskinesia; Huntington's disease; Gilles de la Tourette's syndrome;
OS Homo sapiens.
PN EP-866126-A1.
PD 23-SEP-1998.
PF 16-FEB-1998; 301122.
PR 19-MAR-1997; US-820521.
PA (SMK) SMITHKLINE BEECHAM CORP.
PI Bergsma DJ, Fuetterer WS, Mao UT, Sathe GM;
DR WPI: 98-482962/42.
DR N-PSDB: V53631.
PT New polynucleotides and polypeptides encoding a novel human
PT 7-transmembrane receptor - useful for diagnosing and treating e.g.
PT cancer, osteoporosis and Parkinson's disease and infections caused
PT by HIV-1 or -2.
PS Claim 1; Page 18-19; 24pp; English.
CC This polypeptide comprises HNFY20, a novel human 7-transmembrane
CC G-protein coupled receptor that shows about 30.8% identity in 299
CC amino acid residues with the thrombin receptor. Its amino acid
CC sequence was deduced from an isolated HNFY20 polynucleotide
CC sequence (see V53631). The invention relates to HNFY20
CC polypeptides and recombinant materials and methods for their
CC production. It also provides methods for using such polypeptides
CC and HNFY20 polynucleotides for treatment of infections such as
CC bacterial, fungal, protozoan and particularly HIV-1 or HIV-2
CC infections, and conditions including pain, cancers, anorexia,
CC asthma, Parkinson's disease, acute heart failure, hypertension,
CC hypertension, urinary retention, osteoporosis, angina pectoris,
CC myocardial infarction, ulcers, allergies, benign prostatic
CC hypertrophy, and psychotic and neurological disorders, including
CC anxiety, schizophrenia, manic depression, delirium, dementia,
CC severe mental retardation and dyskinesias, such as Huntington's
CC disease or Gilles de la Tourette's syndrome. Gene therapy using
CC RNA encoding HNFY20 can be used to treat conditions caused by
CC under-expression of the protein. The invention also relates to
CC methods of identifying agonists and antagonists and for using
CC such compounds to treat conditions associated with HNFY20
CC imbalance. Diagnostic assays for detecting diseases associated
CC with inappropriate HNFY20 activity or levels are also provided.
SQ Sequence 401 AA;
Query Match 14.5%; Score 81; DB 35; Length 401;
Best Local Similarity 38.1%; Pred. No. 8.50e+01;
Matches 8; Conservative 9; Mismatches 4; Indels 0; Gaps 0;

Db 332 yfsssgfqaafhellrrlqgl 352
 QY 34 FLINSSYESSFLELLEKLT 54

RESULT 11
 ID W10038 standard; Protein: 1253 AA.
 AC W10038:
 DE 07-FEB-1998 (first entry)
 KW Mad binding protein, mslnA.
 KW murine; mslnA; mammalian homologue; Saccharomyces cerevisiae; repressor;
 KW Sin3; Mad; Max; msln:Mad complex; msln:Mad:Max complex; Myc; promoter;
 KW basic helix-loop-helix zipper protein; compete; DNA-binding;
 KW Myc:Max complex; activate; transcription; gene regulation.
 OS Mus musculus.
 FH Key
 FT Misc_difference 10 Location/Qualifiers
 FT /label= unknown
 FT /note= "encoded by TAG"
 FT Misc_difference 1238
 FT /label= unknown
 FT /note= "encoded by TAG"
 FT US5624818-A.
 PN 29-APR-1997.
 PF 01-JUN-1994; 252966.
 PR 01-JUN-1994; US-252966.
 PR 19-SEP-1991; US-756195.
 PR 23-JUN-1992; US-903710.
 PR 01-APR-1994; US-222638.
 PA (HUTC-) HUTCHINSON CANCER RES CENT FRED.
 PI Ayer DE, Eisenman RN;
 DR WPI: 97-258216/23.
 DR N-PSDB: T70126.
 PT msln nucleic acids encoding recombinant polypeptide(s) that
 PT associate with Mad polypeptide are possible homologues of S.
 PT cerevisiae general repressor protein Sin3
 PS Example 12; Fig 23A-C; 11pp; English.
 CC This sequence represents the murine protein, designated mslnA, which may
 CC be a mammalian homologue of the Saccharomyces cerevisiae general
 CC repressor protein Sin3. The msln protein associates with a Mad
 CC polypeptide to form a msln:Mad complex, which preferably associates
 CC with a Max polypeptide to form a msln:Mad:Max complex which binds to a
 CC nucleotide sequence comprising CACGTG. Mad is a basic helix-loop-helix
 CC (bHLH) zipper protein which can compete with Myc by forming sequence-
 CC specific DNA-binding heterocomplexes with Max. Mad:Max complexes repress,
 CC while Myc:Max complexes activate, transcription from promoters containing
 CC proximal CACGTG binding sites for these proteins. Expression of Mad is
 CC closely linked to differentiation in at least two distinct cell lineages.
 CC The switch from Myc:Max to Mad:Max complexes may reflect the repression
 CC of transcription of Myc regulated genes by Mad. The DNA, vectors and host
 CC cells of the invention are useful for the recombinant production of msln
 CC proteins useful in elucidation of Mad repressor functions.
 SQ Sequence 1253 AA;
 Query Match 14.5%; Score 81; DB 25; Length 1253;
 Best Local Similarity 27.5%; Pred. No. 8.50e+01;
 Matches 11; Conservative 14; Mismatches 14; Indels 1; Gaps 1;
 Db 949 vlgikrdsdpaiqlrlkpemdvdydyafidmrsl 988
 QY 13 LTGSSHGTPGWTQLKIKESF-LTNSSYESSFLELLEKLT 51
 RESULT 12
 ID W10040 standard; Protein: 1261 AA.
 AC W10040:
 DE 07-FEB-1998 (first entry)
 KW Mad binding protein, mslnA9.
 KW murine; mslnA; mammalian homologue; Saccharomyces cerevisiae; repressor;
 KW Sin3; Mad; Max; msln:Mad complex; msln:Mad:Max complex; Myc; promoter;
 KW basic helix-loop-helix zipper protein; compete; DNA-binding;
 KW Myc:Max complex; activate; transcription; gene regulation.

OS Mus musculus.
 FH Key
 FT Misc_difference 10 Location/Qualifiers
 FT /label= unknown
 FT /note= "encoded by TAG"
 FT Misc_difference 1247
 FT /label= unknown
 FT /note= "encoded by TAG"
 FT US5624818-A.
 PN 29-APR-1997.
 PF 01-JUN-1994; 252966.
 PR 01-JUN-1994; US-252966.
 PR 19-SEP-1991; US-756195.
 PR 23-JUN-1992; US-903710.
 PR 01-APR-1994; US-222638.
 PA (HUTC-) HUTCHINSON CANCER RES CENT FRED.
 PI Ayer DE, Eisenman RN;
 DR WPI: 97-258216/23.
 DR N-PSDB: T70128.
 PT msln nucleic acids encoding recombinant polypeptide(s) that
 PT associate with Mad polypeptide - are possible homologues of S.
 PT cerevisiae general repressor protein Sin3
 PS Example 12; Fig 29A-C; 11pp; English.
 CC This sequence represents the murine protein, designated mslnA9, which
 CC may be a mammalian homologue of the Saccharomyces cerevisiae general
 CC repressor protein Sin3. The msln protein associates with a Mad
 CC polypeptide to form a msln:Mad complex, which preferably associates
 CC with a Max polypeptide to form a msln:Mad:Max complex which binds to a
 CC nucleotide sequence comprising CACGTG. Mad is a basic helix-loop-helix
 CC (bHLH) zipper protein which can compete with Myc by forming sequence-
 CC specific DNA-binding heterocomplexes with Max. Mad:Max complexes repress,
 CC while Myc:Max complexes activate, transcription from promoters containing
 CC proximal CACGTG binding sites for these proteins. Expression of Mad is
 CC closely linked to differentiation in at least two distinct cell lineages.
 CC The switch from Myc:Max to Mad:Max complexes may reflect the repression
 CC of transcription of Myc regulated genes by Mad. The DNA, vectors and host
 CC cells of the invention are useful for the recombinant production of msln
 CC proteins useful in elucidation of Mad repressor functions.
 SQ Sequence 1261 AA;
 Query Match 14.5%; Score 81; DB 25; Length 1261;
 Best Local Similarity 27.5%; Pred. No. 8.50e+01;
 Matches 11; Conservative 14; Mismatches 14; Indels 1; Gaps 1;
 Db 949 vlgikrdsdpaiqlrlkpemdvdydyafidmrsl 988
 QY 13 LTGSSHGTPGWTQLKIKESF-LTNSSYESSFLELLEKLT 51
 RESULT 13
 ID R15138 standard; Protein: 398 AA.
 AC R15138:
 DE 17-FEB-1992 (first entry)
 DE Human serotonin 1D receptor encoded by gene 11.
 KW 5-HT (1D); Parkinson's Disease; migraine; anxiety; eating disorder;
 KW G-protein; 5-hydroxytryptamine.
 OS Homo sapiens.
 FH Key
 FT domain Location/Qualifiers
 FT 1..49 /label= extracellular
 FT 50..73 /label= transmembrane-1
 FT 74..86 /label= transmembrane-1
 FT 87..110 /label= cytoplasmic
 FT 111..121 /label= transmembrane-2
 FT 122..145 /label= extracellular
 FT 146..164 /label= transmembrane-3
 FT 165..188 /label= cytoplasmic
 FT domain

[M] [P] [E] [R] [E] [I] [T] (TM)

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MPerch_app protein - protein database search, using Smith-Waterman algorithm

Run on: Fri Oct 22 18:45:15 1999, MasPar time 2.38 Seconds

Tabular output not generated. 383.184 Million cell updates/sec

Title: >US-09-092-296-15
Description: (1-78) from US09092296.pep
Perfect Score: 558
Sequence: 1 MGSLPLVLLILILGSSHGT.....SGTSTVLHARSQHVVCNT 78

Scoring table: PAM 150
Gap 11

Searched: 119857 seqs, 11713122 residues

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database:

a-issued
1:5A_COMB 2:5B_COMB 3:PCT9_COMB 4:backfiles1

Statistics: Mean 25.509; Variance 122.762; scale 0.208

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description	Pred. No.
1	85	15.2	325	1	US-08-118-	Sequence 51, Applicati	2.00e+01
2	85	15.2	325	3	PCT-US93-0	Sequence 51, Applicati	2.00e+01
3	83	14.9	194	1	US-08-063-	Sequence 8, Applicatio	2.75e+01
4	83	14.9	194	3	PCT-US93-0	Sequence 8, Applicatio	2.75e+01
5	82	14.7	169	1	US-08-741-	Sequence 2, Applicatio	3.23e+01
6	81	14.5	401	2	US-08-820-	Sequence 2, Applicatio	3.78e+01
7	81	14.5	1253	1	US-08-252-	Sequence 12, Applicati	3.78e+01
8	81	14.5	1261	1	US-08-252-	Sequence 18, Applicati	3.78e+01
9	80	14.3	169	2	US-08-828-	Sequence 6, Applicatio	4.43e+01
10	80	14.3	390	1	US-07-817-	Sequence 6, Applicatio	4.43e+01
11	80	14.3	390	1	US-08-117-	Sequence 6, Applicatio	4.43e+01
12	80	14.3	390	3	PCT-US93-0	Sequence 6, Applicatio	4.43e+01
13	80	14.3	390	1	US-08-216-	Sequence 6, Applicatio	4.43e+01
14	80	14.3	398	2	US-08-542-	Sequence 6, Applicatio	4.43e+01
15	80	14.3	398	1	US-08-310-	Sequence 6, Applicatio	4.43e+01
16	80	14.3	1214	2	US-08-231-	Sequence 54, Applicati	4.43e+01
17	80	14.3	1219	2	US-08-231-	Sequence 50, Applicati	4.43e+01
18	80	14.3	1231	2	US-08-231-	Sequence 48, Applicati	4.43e+01
19	80	14.3	1235	2	US-08-231-	Sequence 6, Applicatio	4.43e+01
20	80	14.3	1239	2	US-08-231-	Sequence 52, Applicati	4.43e+01
21	80	14.3	1244	2	US-08-231-	Sequence 46, Applicati	4.43e+01
22	79	14.2	1229	2	US-08-339-	Sequence 33, Applicati	5.18e+01
23	78	14.0	120	2	US-08-420-	Sequence 29, Applicati	6.06e+01

24	78	14.0	120	3	PCT-US93-1	Sequence 29, Applicati	6.06e+01
25	78	14.0	1239	1	US-08-026-	Sequence 3, Applicatio	6.06e+01
26	76	13.6	197	3	PCT-US93-0	Sequence 2, Applicatio	8.28e+01
27	76	13.6	197	2	US-08-215-	Sequence 2, Applicatio	8.28e+01
28	76	13.6	317	2	US-09-213-	Sequence 3, Applicatio	8.28e+01
29	76	13.6	318	2	US-08-619-	Sequence 3, Applicatio	8.28e+01
30	76	13.6	340	2	US-09-213-	Sequence 1, Applicatio	8.28e+01
31	76	13.6	477	4	US-09-213-	Sequence 1, Applicatio	8.28e+01
32	76	13.6	477	1	5245013-3	Patent No. 5245013.	8.28e+01
33	76	13.6	477	1	US-07-847-	Sequence 2, Applicatio	8.28e+01
34	76	13.6	477	2	US-08-990-	Sequence 2, Applicatio	8.28e+01
35	76	13.6	481	1	US-08-306-	Sequence 98, Applicati	8.28e+01
36	76	13.6	481	1	US-08-261-	Sequence 4, Applicatio	8.28e+01
37	76	13.6	481	2	US-08-274-	Sequence 5, Applicatio	8.28e+01
38	76	13.6	481	2	US-08-186-	Sequence 2, Applicatio	8.28e+01
39	76	13.6	481	1	US-08-485-	Sequence 98, Applicati	8.28e+01
40	76	13.6	481	3	PCT-US93-0	Sequence 9, Applicatio	8.28e+01
41	76	13.6	481	3	PCT-US93-0	Sequence 98, Applicati	8.28e+01
42	76	13.6	481	3	PCT-US94-0	Sequence 98, Applicati	8.28e+01
43	76	13.6	481	2	US-08-215-	Sequence 98, Applicati	8.28e+01
44	76	13.6	481	3	PCT-US94-0	Sequence 4, Applicatio	8.28e+01
45	76	13.6	528	2	US-08-403-	Sequence 21, Applicati	8.28e+01

ALIGNMENTS

RESULT 1
ID US-08-118-270-51 STANDARD: PRT; 325 AA.

AC xxxxxx

DE Sequence 51, Application US/08118270

CC Patent No. 5508384

CC GENERAL INFORMATION:

CC APPLICANT: Murphy, Randall B.

CC ATTORNEY/AGENT INFORMATION:

CC TITLE OF INVENTION: POLYPEPTIDES OF G-COUPLED PROTEIN

CC RECEPTORS, AND COMPOSITIONS AND METHODS THEREOF

CC NUMBER OF SEQUENCES: 348

CC CORRESPONDENCE ADDRESS:

CC ADDRESSEE: BROWDY AND NEWMARK

CC STREET: 419 Seventh Street, N.W., Suite 300

CC CITY: Washington

CC STATE: D.C.

CC COUNTRY: USA

CC ZIP: 20004

CC COMPUTER READABLE FORM:

CC MEDIUM TYPE: Floppy disk

CC COMPUTER: IBM PC compatible

CC OPERATING SYSTEM: PC-DOS/MS-DOS

CC SOFTWARE: Patent Release #1.0, Version #1.25

CC CURRENT APPLICATION DATA:

CC APPLICATION NUMBER: US/08/118,270

CC FILING DATE: 09-SEP-1993

CC PRIORITY APPLICATION DATA:

CC APPLICATION NUMBER: US 07/943,236

CC FILING DATE: 10-SEP-1992

CC ATTORNEY/AGENT INFORMATION:

CC NAME: Townsend, Kevin G.

CC REGISTRATION NUMBER: 34,033

CC TELEPHONE/DOCKET NUMBER: MURPHY-2A

CC TELEPHONE: 202-628-5197

CC TELEFAX: 202-737-5528

CC TELEX: 248633

CC INFORMATION FOR SEQ ID NO: 51:

CC SEQUENCE CHARACTERISTICS:

CC LENGTH: 325 amino acids

CC TYPE: amino acid

CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: peptide
SQ SEQUENCE 325 AA; 3728 MM; 592694 CN;

Query Match 15.2%; Score 85; DB 1; Length 325;
Best Local Similarity 44.4%; Pred. No. 2.00e+01;
Matches 12; Conservative 5; Mismatches 8; Indels 2;

Db 27 YMKLKYASVFLNLADLCFLITLP 53
QY 34 FLTNSYESSFL-EL-LEKLCILLHLP 58

RESULT 2
ID PCT-US93-08528-51 STANDARD; PRT: 325 AA.
XX
XX
XX xxxxxx

Sequence 51, Application PC/TUS9308528

Sequence 51, Application PC/TUS9308528

GENERAL INFORMATION:

APPLICANT: New York University

TITLE OF INVENTION: POLYPEPTIDES OF G-COUPLED PROTEIN

NUMBER OF INVENTION: RECEPTORS, AND COMPOSITIONS AND METHODS THEREOF

NUMBER OF SEQUENCES: 348

CORRESPONDENCE ADDRESS:

ADDRESSEE: BROMDY AND NEIMARK

STREET: 419 Seventh Street, N.W., Suite 300

CITY: Washington

STATE: D.C.

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: PCT/US93/08528

FILING DATE: 09-SEP-1993

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/943,236

FILING DATE: 10-SEP-1992

ATTORNEY/AGENT INFORMATION:

NAME: Townsend, Kevin G.

REGISTRATION NUMBER: 34,033

REFERENCE/DOCKET NUMBER: WURPHY-2 PCT

TELECOMMUNICATION INFORMATION:

TELEPHONE: 202-628-5197

TELEFAX: 202-737-3528

TELEX: 248633

INFORMATION FOR SEQ ID NO: 51:

SEQUENCE CHARACTERISTICS:

LENGTH: 325 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: peptide

SEQUENCE 325 AA; 3728 MM; 592694 CN;

Query Match 15.2%; Score 85; DB 3; Length 325;
Best Local Similarity 44.4%; Pred. No. 2.00e+01;
Matches 12; Conservative 5; Mismatches 8; Indels 2; Gaps 2;

Db 27 YMKLKYASVFLNLADLCFLITLP 53
QY 34 FLTNSYESSFL-EL-LEKLCILLHLP 58

RESULT 3
ID US-08-063-552-8 STANDARD; PRT: 194 AA.
XX
XX
XX xxxxxx

Sequence 8, Application US/08063552

Db 27 YMKLKYASVFLNLADLCFLITLP 53
QY 34 FLTNSYESSFL-EL-LEKLCILLHLP 58

Sequence 8, Application US/08063552

Sequence 8, Application US/08063552

Sequence 8, Application US/08063552

Sequence 8, Application US/08063552

GENERAL INFORMATION:

APPLICANT: Edwards, Robert H

TITLE OF INVENTION: Vesicle Membrane Transport Proteins

NUMBER OF SEQUENCES: 17

CORRESPONDENCE ADDRESS:

ADDRESSEE: Sheldon & Max

STREET: 225 South Lake Avenue, Ninth Floor

CITY: Pasadena

STATE: California

COUNTRY: USA

ZIP: 91101

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/063,552

FILING DATE: 19930514

CLASSIFICATION: 530

ATTORNEY/AGENT INFORMATION:

NAME: Farber, Michael B

REGISTRATION NUMBER: 32,612

REFERENCE/DOCKET NUMBER: 9067-1

TELECOMMUNICATION INFORMATION:

TELEPHONE: (818) 796-4000

TELEFAX: (818) 795-6321

INFORMATION FOR SEQ ID NO: 8:

SEQUENCE CHARACTERISTICS:

LENGTH: 194 amino acids

TYPE: AMINO ACID

TOPOLOGY: linear

MOLECULE TYPE: peptide

HYPOTHETICAL: NO

FRAGMENT TYPE: Internal

ORIGINAL SOURCE:

ORGANISM: Transposon 10

SEQUENCE 194 AA; 20868 MM; 201442 CN;

Query Match 14.9%; Score 83; DB 1; Length 194;
Best Local Similarity 31.0%; Pred. No. 2.75e+01;
Matches 18; Conservative 13; Mismatches 23; Indels 4; Gaps 4;

Db 1 MNSRTIAVITLIDAM-GIGLIMVLPILARE-FIASDIDNHGVULA-LVALMOV 55
QY 1 MGSGLPVLVLTLLGSSHGTPGPM-LQLKLKESFLTNSYSFLELVLEKLCILLHLP 57

RESULT 4
ID PCT-US93-05704-8 STANDARD; PRT: 194 AA.
XX
XX
XX xxxxxx

Sequence 8, Application PC/TUS9305704

Sequence 8, Application PC/TUS9305704

GENERAL INFORMATION:

APPLICANT: Edwards, Robert H

TITLE OF INVENTION: Vesicle Membrane Transport Proteins

NUMBER OF SEQUENCES: 17

CC CORRESPONDENCE ADDRESS:
 CC ADDRESSEE: Sheldon & Max
 CC STREET: 225 South Lake Avenue, Ninth Floor
 CC CITY: Pasadena
 CC STATE: California
 CC COUNTRY: USA
 CC ZIP: 91001
 CC COMPUTER READABLE FORM:
 CC MEDIUM TYPE: Floppy disk
 CC COMPUTER: IBM PC compatible
 CC OPERATING SYSTEM: PC-DOS/MS-DOS
 CC SOFTWARE: Patent In Release #1.0, Version #1.25
 CC CURRENT APPLICATION DATA:
 CC APPLICATION NUMBER: PCT/US93/05704
 CC FILING DATE: 19930611
 CC CLASSIFICATION:
 CC ATTORNEY/AGENT INFORMATION:
 CC NAME: Faider, Michael B
 CC REGISTRATION NUMBER: 32,612
 CC REFERENCE/DOCKET NUMBER: 9067-1PCT
 CC TELECOMMUNICATION INFORMATION:
 CC TELEPHONE: (818) 796-4000
 CC TELEFAX: (818) 795-6321
 CC INFORMATION FOR SEQ ID NO: 8:
 CC SEQUENCE CHARACTERISTICS:
 CC LENGTH: 194 amino acids
 CC TYPE: AMINO ACID
 CC TOPOLOGY: linear
 CC MOLECULE TYPE: Peptide
 CC HYPOTHEICAL: NO
 CC FRAGMENT TYPE: Internal
 CC ORIGINAL SOURCE:
 CC ORGANISM: transposon 10
 CC SEQUENCE 194 AA; 20868 MW; 201442 CN;
 SQ
 Query Match 14.9%; Score 83; DB 3; Length 194;
 Best Local Similarity 31.0%; Pred. No. 2.75e+01;
 Matches 18; Conservative 13; Mismatches 23; Indels 4; Gaps 4;
 DB 1 MNSSTIALVITLDAM-GIGLIMPVPLRLRE-PIASEDIANHEGVLLA-LYALMOV 55
 QY 1 MGSGLPVLTLTGLSSHGTPGMT-LQIKRESPLTNSSESSFELLEKICLLHL 57
 RESULT 5
 ID US-08-741-406-2 STANDARD; PRT; 169 AA.
 AC xxxxxx
 DE
 XX
 XX
 DT
 DT
 DE
 XX
 XX
 CC Sequence 2, Application US/08741406
 CC Patent No. 5721118
 CC GENERAL INFORMATION:
 CC APPLICANT: Schettler, Immo E.
 CC TITLE OF INVENTION: Mammalian Artificial Chromosomes and
 CC TITLE OF INVENTION: Methods of Using Same
 CC NUMBER OF SEQUENCES: 16
 CC CORRESPONDENCE ADDRESS:
 CC ADDRESSEE: Campbell & Flores LLP
 CC STREET: 4370 La Jolla Village Drive, Suite 700
 CC CITY: San Diego
 CC STATE: California
 CC COUNTRY: United States
 CC ZIP: 92122
 CC COMPUTER READABLE FORM:
 CC MEDIUM TYPE: Floppy disk
 CC COMPUTER: IBM PC compatible
 CC OPERATING SYSTEM: PC-DOS/MS-DOS
 CC SOFTWARE: Patent In Release #1.0, Version #1.25
 CC CURRENT APPLICATION DATA:

CC APPLICATION NUMBER: US/08/741,406
 CC FILING DATE:
 CC CLASSIFICATION: 514
 CC PRIOR APPLICATION DATA:
 CC APPLICATION NUMBER: US 08/550,717
 CC FILING DATE: 31-OCT-1995
 CC ATTORNEY/AGENT INFORMATION:
 CC NAME: Campbell, Cathryn A.
 CC REGISTRATION NUMBER: 31,815
 CC REFERENCE/DOCKET NUMBER: P-UD 2317
 CC TELECOMMUNICATION INFORMATION:
 CC TELEPHONE: (619) 535-9001
 CC TELEFAX: (619) 535-8949
 CC INFORMATION FOR SEQ ID NO: 2:
 CC SEQUENCE CHARACTERISTICS:
 CC LENGTH: 169 amino acids
 CC TYPE: amino acid
 CC TOPOLOGY: linear
 CC MOLECULE TYPE: protein
 CC SEQUENCE 169 AA; 18610 MW; 162524 CN;
 SQ
 Query Match 14.7%; Score 82; DB 1; Length 169;
 Best Local Similarity 28.0%; Pred. No. 3.23e+01;
 Matches 14; Conservative 20; Mismatches 13; Indels 3; Gaps 3;
 DB 62 SLPMASICHRTGAINASGVSL-EGMS-ALLPBNFES-YIELYKSICL 108
 QY 4 GLPVLTLTGLSSHGTPGMTLQIKRESPLTNSSESSFELLEKICL 53
 RESULT 6
 ID US-08-820-521-2 STANDARD; PRT; 401 AA.
 AC xxxxxx
 DE
 XX
 XX
 DT
 DT
 DE
 XX
 XX
 CC Sequence 2, Application US/08820521
 CC Patent No. 5942416
 CC GENERAL INFORMATION:
 CC APPLICANT: Bergema, Derk
 CC APPLICANT: Ganesu, Sathe
 CC APPLICANT: Fuetterer, Wendy
 CC APPLICANT: Mao, Joyce
 CC TITLE OF INVENTION: CDNA CLONE HNFY20 THAT ENCODES
 CC TITLE OF INVENTION: A NOVEL HUMAN 7-TRANSMEMBRANE RECEPTOR
 CC NUMBER OF SEQUENCES: 2
 CC CORRESPONDENCE ADDRESS:
 CC ADDRESSEE: Smithline Beecham Corporation
 CC STREET: 709 Swedeland Road
 CC CITY: King of Prussia
 CC STATE: PA
 CC COUNTRY: USA
 CC ZIP: 19406
 CC COMPUTER READABLE FORM:
 CC MEDIUM TYPE: Diskette
 CC COMPUTER: IBM Compatible
 CC OPERATING SYSTEM: DOS
 CC SOFTWARE: FASTSEQ for Windows Version 2.0
 CC CURRENT APPLICATION DATA:
 CC APPLICATION NUMBER: US/08/820,521
 CC FILING DATE: 19-MAR-1997
 CC CLASSIFICATION: 435
 CC PRIOR APPLICATION DATA:
 CC APPLICATION NUMBER:
 CC FILING DATE:
 CC ATTORNEY/AGENT INFORMATION:
 CC NAME: Han, William T
 CC REGISTRATION NUMBER: 34,344
 CC REFERENCE/DOCKET NUMBER: GH50011
 CC TELECOMMUNICATION INFORMATION:

```

CC      TELEPHONE: 610-270-5219
CC      TELEFAX: 610-270-4026
CC      TELEX:
CC      INFORMATION FOR SEQ ID NO: 2:
CC      SEQUENCE CHARACTERISTICS:
CC          LENGTH: 401 amino acids
CC          TYPE: amino acid
CC          STRANDEDNESS: single
CC          TOPOLOGY: linear
CC      MOLECULE TYPE: protein
CC      SEQUENCE 401 AA: 44386 MW: 853771 CN:
SQ
Query Match          14.5%  Score 81;  DB 2;  Length 401;
Best Local Similarity 38.1%;  Pred. No. 3,78e+01;
Matches      8;  Conservative      9;  Mismatches      4;  Indels      0;  Gaps      0;
0y.      34  YFSSSGQADPELLLRDGL 352
          :::::::::::|::|::|::|
XX      US-08-252-966B-12      STANDARD:      PRT;      1253 AA.
XX      AC      xxxxxx
XX      DT
XX
DE      Sequence 12, Application US/08252966B
XX
XX      Sequence 12, Application US/08252966B
XX      Patent No. 5624818
CC      GENERAL INFORMATION:
CC      APPLICANT: EISENMAN, Robert N.
CC      APPLICANT: HURLIN, Peter J.
CC      APPLICANT: AYER, Donald E.
CC      TITLE OF INVENTION: Regulatory Proteins that Dimerize with
CC      TITLE OF INVENTION: Mad or Max
CC      NUMBER OF SEQUENCES: 19
CC
CC      CORRESPONDENCE ADDRESS:
CC      ADDRESSEE: Christensen, O'Connor, Johnson, and KindnessP/LC
CC      STREET: 1420 Fifth Ave., Suite 2800
CC      CITY: Seattle
CC      STATE: Washington
CC      COUNTRY: USA
CC      ZIP: 98101-2347
CC      COMPUTER READABLE FORM:
CC      MEDIUM TYPE: Floppy disk
CC      COMPUTER: IBM PC compatible
CC      OPERATING SYSTEM: PC-DOS/MS-DOS
CC      SOFTWARE: Patentin Release #1.0, Version #1.25
CC      CURRENT APPLICATION DATA:
CC      APPLICATION NUMBER: US/08/252,966B
CC      FILING DATE: 01-JUN-1994
CC      CLASSIFICATION: 435
CC      ATTORNEY/AGENT INFORMATION:
CC      NAME: Shelton, Dennis K.
CC      REGISTRATION NUMBER: 26,997.
CC      REFERENCE/DOCKET NUMBER: PHC17694
CC      TELECOMMUNICATION INFORMATION:
CC      TELEPHONE: (206) 682-8100
CC      TELEFAX: (206) 224-0779
CC      INFORMATION FOR SEQ ID NO: 12:
CC      SEQUENCE CHARACTERISTICS:
CC          LENGTH: 1253 amino acids
CC          TYPE: amino acid
CC          STRANDEDNESS: single
CC          TOPOLOGY: linear
CC      MOLECULE TYPE: protein
CC      DESCRIPTION: translation of msina cDNA; see Figure 23
CC      HYDROPHILIC: YES
CC      ORIGINAL SOURCE:
CC      ORGANISM: Mus musculus

```

```

SQ SEQUENCE 1253 AA: 142589 MW: 7863283 CN:
Query Match . 14.5% Score 81; DB 1; Length 1253;
Best Local Similarity 27.5%; Pred. No. 3.78e+01;
Matches 11; Conservative 14; Mismatches 14; Indels 1; Gaps 1;
Db 949 VLGIRKSDSPAIQLAKPMQVDEYYPALDMVMSL 988
:11: : : : : : : : : : : : : : : : : : : :
QY 13 LIGSSHGTCGATLQLEKESF-LTNSYESSFLELEKL 51

RESULT 8 STANDARD; PRT; 1261 AA.
XX ID US-08-252-966B-18
XX AC XXXXXX
XX DN
XX DE
XX SE Sequence 18, Application US/08252966B
CC CC Sequence 18, Application US/08252966B
CC CC Patent No. 5624818
CC CC GENERAL INFORMATION:
CC CC APPLICANT: Eismann, Robert N.
CC CC APPLICANT: Hurlin, Peter J.
CC CC APPLICANT: Ayer, Donald E.
CC CC TITLE OF INVENTION: Regulatory Proteins that Dimerize with
CC CC TITLE OF INVENTION: Mad or Max
CC CC NUMBER OF SEQUENCES: 19
CC CC CORRESPONDENCE ADDRESS:
CC CC ADDRESSEE: Christensen, O'Connor, Johnson, and KindnessPLLC
CC CC STREET: 1420 Fifth Ave., Suite 2800
CC CC CITY: Seattle
CC CC STATE: Washington
CC CC COUNTRY: USA
CC CC ZIP: 98101-2347
CC CC COMPUTER READABLE FORM:
CC CC MEDIUM TYPE: Floppy disk
CC CC COMPUTER: IBM PC compatible
CC CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC CC SOFTWARE: Patentin Release #1.0, Version #1.25
CC CC CURRENT APPLICATION DATA:
CC CC APPLICATION NUMBER: US/08/252,966B
CC CC FILING DATE: 01-JUN-1994
CC CC CLASSIFICATION: 435
CC CC ATTORNEY/AGENT INFORMATION:
CC CC NAME: Shelton, Dennis K.
CC CC REGISTRATION NUMBER: 26,997.
CC CC REFERENCE/DOCKET NUMBER: FHC817694
CC CC TELECOMMUNICATION INFORMATION:
CC CC TELEPHONE: (206) 682-8100
CC CC TELEFAX: (206) 224-0779
CC CC INFORMATION FOR SEQ ID NO: 18:
CC CC SEQUENCE CHARACTERISTICS:
CC CC LENGTH: 1261 amino acids
CC CC TYPE: amino acid
CC CC STRANDEDNESS: single
CC CC TOPOLOGY: linear
CC CC MOLECULE TYPE: protein
CC CC DESCRIPTION: translation of msina9 cDNA; see Figure 29A, B, C, D
CC CC HYPOTHEetical: YES
CC CC ORIGINAL SOURCE:
CC CC ORGANISM: Mus musculus
CC CC SEQUENCE 1261 AA; 143711 MW; 7937040 CN;

Query Match 14.5% Score 81; DB 1; Length 1261;
Best Local Similarity 27.5%; Pred. No. 3.78e+01;
Matches 11; Conservative 14; Mismatches 14; Indels 1; Gaps 1
Ddb 949 VLGIRKSDSPAIQLAKPMQVDEYYPALDMVMSL 988
:11: : : : : : : : : : : : : : : : : : : :
QY 13 LIGSSHGTCGATLQLEKESF-LTNSYESSFLELEKL 51

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[illegible]

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CC Sequence 6, Application US/07817920
CC Patent No. 5360735
CC GENERAL INFORMATION:
CC APPLICANT: Weinshank, Richard L
CC APPLICANT: Branchak, Theresa
CC APPLICANT: Hatcliff, Paul R
CC TITLE OF INVENTION: DNA ENCODING A HUMAN 5-HT1F RECEPTOR AND
CC TITLE OF INVENTION: US5 THEREOF
CC NUMBER OF SEQUENCES: 9
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Cooper & Dunham
CC STREET: 30 Rockefeller Plaza
CC CITY: New York
CC STATE: New York
CC COUNTRY: USA
CC ZIP: 10112
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patent in Release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/07/817,920
CC FILING DATE: 19920108
CC CLASSIFICATION: 514
CC ATTORNEY/AGENT INFORMATION:
CC NAME: White, John P
CC REGISTRATION NUMBER: 28,678
CC REFERENCE/DOCKET NUMBER: 1795/39318
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: 212-977-9950
CC TELEFAX: 212-664-0525
CC CC TELEX: 422523 COOP UT
CC INFORMATION FOR SEQ ID NO: 6:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 390 amino acids
CC TYPE: AMINO ACID
CC STRANDEDNESS: unknown
CC TOPOLOGY: linear
CC MOLECULE TYPE: protein
CC HYPOTHEICAL: NO
CC ANTI-SENSE: NO
CC FRAGMENT TYPE: N-terminal
CC IMMEDIATE SOURCE:
CC CLONE: 5-HT1DB
CC CC SEQUENCE 390 AA; 43656 MW; 849817 CN;
SQ
Query Match 14.23; Score 80; DB 1; Length 390.
Best Local Similarity 23.65; Fred. No. 4.43e+01;
Matches 17; Conservative 22; Mismatches 32; Indels 1; Gaps 1.
Db
52 LVMLLALITLATTTSSNAFVATVYRTRRLTPANTPLNLSIDPVTDLVSLIVIPISMTYV 111
|||::: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 7 LVLLLLILSSIGTGPMILDLKLKESFL-TNSYSSFFFLIKELCLLHLPGSGSVTL 65
:: : : : :
Db 112 TDRWTLISOVCD 123
:: : : : :
QY 66 HHARSQHYYCN 77
RESULT 11
ID US-08-117-006-6 STANDARD; PRT, 390 AA.
XX
AC xxxxxx
DT
DE Sequence 6, Application US/08117006
CC Sequence 6, Application US/08117006
CC Patent No. 5639652
CC GENERAL INFORMATION:
CC APPLICANT: Weinshank, Richard L.
CC

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CC      PRIOR APPLICATION DATA:
CC      APPLICATION NUMBER: US/08/194,113
CC      FILING DATE:
CC      APPLICATION NUMBER: US/07/803,626
CC      FILING DATE:
CC      ATTORNEY/AGENT INFORMATION:
CC      NAME: White, John P.
CC      REGISTRATION NUMBER: 28,678
CC      REFERENCE/DOCKET NUMBER: 1795/39317
CC      TELECOMMUNICATION INFORMATION:
CC      TELEPHONE: 212-977-9550
CC      TELEFAX: 212-664-0525
CC      TELEX: 422523 COOP U1
CC      INFORMATION FOR SEQ ID NO: 6:
CC      SEQUENCE CHARACTERISTICS:
CC      LENGTH: 398 amino acids
CC      TYPE: amino acid
CC      STRANDEDNESS: unknown
CC      TOPOLOGY: unknown
CC      MOLECULE TYPE: protein
CC      HYPOTHEetical: NO
CC      ANTI-SENSE: NO
CC      FRAGMENT TYPE: N-terminal
SO      SEQUENCE 398 AA: 44384 MW: 880684 CN:

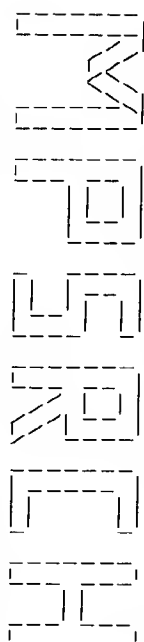
Query Match      14.3%  Score 80: DB 1: Length 398:
Best Local Similarity 23.5%  Pired. No. 4,438+01;
Matches 17: Conservative 22; Mismatches 32; Indels 1; Caps 1;

Db 52 LVMALLATLTLNAFVATVYRTKLTNPANYLIASLDVTDLLVSILVPISTMTV 111
Oy 7 LVLLLLSSHGTPGPTQLKLKESFL-TNSYESSFLELKLCLLLHLPSTSVTL 65

Db 112 TDRRTLSQVYCD 123
Oy 66 HHRASQHVYCN 77

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Search completed: Fri Oct 22 18:45:24 1999
Job time : 9 secs.



(TM)

Release 3.1a John F. Collins, Biocomputing Research Unit.
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Distribution rights by Oxford Molecular Ltd

MSrch_PP protein - protein database search, using Smith-Waterman algorithm
Run on: Fri Oct 22 18:42:40 1999; Maspar time 5.89 Seconds
Tabular output not generated. 530.469 Million cell updates/sec

Title: >US-09-092-296-15
Description: (1-78) from US09092296.Pep
Perfect Score: 558
Sequence: 1 MSGGLEPLVLLTLIGSSHGR.....SGTSVTLHARQSHVVCNT 78

Scoring table:
PAM 150
Gap 11

Searched: 122810 seqs, 4006893 residues

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database: pIR60
1:pir1 2:pir2 3:pir3 4:pir4

Statistics: Mean 39.113; Variance 85.189; scale 0.459

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description	Pred. No.
1	101	18.1	501	2	protein export membra	7.90e-02
2	93	16.7	382	2	ubiquinol--cytochrome	8.03e-01
3	92	16.5	712	2	translational regulator	1.07e+00
4	91	16.3	252	2	osmotic-like protein	1.41e+00
5	89	15.9	879	2	hypothetical protein	2.45e+00
6	89	15.9	3005	2	hemocytic protein zfh	2.45e+00
7	88	15.8	1822	2	K12H.8 protein - Cae	3.23e+00
8	87	15.6	143	2	hypothetical protein	4.25e+00
9	87	15.6	396	2	probable translation	5.57e+00
10	86	15.4	272	2	flagellar motor appar	7.28e+00
11	85	15.2	315	2	hypothetical protein	9.51e+00
12	84	15.1	441	2	probable tyrosine kin	9.51e+00
13	84	15.1	454	2	hypothetical protein	1.24e+01
14	84	15.1	509	2	tetracycline resistanc	1.24e+01
15	83	14.9	378	2	beta-galactosidase (E	1.24e+01
16	83	14.9	401	1	mitotic-specific cycl	1.61e+01
17	83	14.9	469	2	T-cell receptor alpha	1.61e+01
18	83	14.9	647	2	T-cell antigen recept	1.61e+01
19	82	14.7	56	2		
20	82	14.7	110	2		
21	82	14.7	110	2		
22	82	14.7	152	2		
23	82	14.7	155	2		

24	82	14.7	372	2	A46138	Invasion protein Inve	1.61e+01
25	82	14.7	946	3	T00024	ent-Kaurene synthase	1.61e+01
26	81	14.5	253	2	S70170	meta-Protein - Rhodop	2.09e+01
27	81	14.5	261	2	I52518	sperm acrosome antigen	2.09e+01
28	81	14.5	287	2	S71192	mitosis-specific cycl	2.09e+01
29	81	14.5	346	2	JC5715	G protein-coupled rec	2.09e+01
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33	81	14.5	482	2	B35843	hypothetical protein	2.09e+01
34	81	14.5	1219	2	I61713	lipopolysaccharide-bi	2.09e+01
35	81	14.5	1229	2	A55068	co-repressor protein	2.09e+01
36	80	14.3	1332	2	A45159	succinate dehydrogena	2.70e+01
37	80	14.3	141	2	G71079	hypothetical protein	2.70e+01
38	80	14.3	191	2	H71370	hypothetical protein	2.70e+01
39	80	14.3	331	2	D70431	oligopeptide transpor	2.70e+01
40	80	14.3	387	1	CBASN	ubiquinol--cytochrome	2.70e+01
41	80	14.3	402	2	S23860	chloramphenicol resis	2.70e+01
42	80	14.3	741	2	I48694	probable transcriptio	2.70e+01
43	80	14.3	742	2	A49672	transcription factor	2.70e+01
44	80	14.3	772	2	A55004	transcription factor	2.70e+01
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ALIGNMENTS

RESULT 1
ENTRY 1 C70384 #type complete
TITLE protein export membrane protein SecD - Aquifex aeolicus
ORGANISM #formal_name Aquifex aeolicus
DATE 08-May-1998 #sequence_revision 08-May-1998 #text_change 12-Feb-1999

ACCESSIONS
#authors A70384
#accession C70384
#status preliminary; nucleic acid sequence not shown; translation not shown
#cross-references MIM:98196666
#accession C70384
#journal #title The complete genome of the hyperthermophilic bacterium Aquifex aeolicus.
Nature (1998) 392:353-358
J.M.; Olson, G.J.; Swanson, R.V.
Keller, M.; Aubay, M.; Huber, R.; Feldman, R.A.; Short, Lenox, A.L.; Graham, D.E.; Overbeck, R.; Sneed, M.A.;
Decker, G.; Warren, P.V.; Gaasterland, T.; Young, W.G.;

GENETICS
#molecule_type DNA
#residues 1-501 #label AOF
#cross-references GB:AE000715; NID:92983478; PID:92983481; GB:AE000657
#experimental_source strain VF5

CLASSIFICATION
#gene secD
#superfamily protein export membrane protein secD
#length 501 #molecular_weight 55459 #checksum 2250

Query Match 18.1%; Score 101; DB 2; Length 501;
Best Local Similarity 38.3%; Pred. No. 7.90e-02;

Matches 18; Conservative 10; Mismatches 18; Indels 1; Gaps 1;
Db 454 VILPFG-GSGPVAGFTATAGTASTSVYAKVFDLINSKITL 499
OY 8 VLLTLGLSSHGTPMDLKLKESFLTNSYSEFFLEKICLL 54

RESULT 2
ENTRY 2 S63638 #type complete
TITLE ubiquinol--cytochrome c reductase (EC 1.10.2.2) cytochrome b
ALTERNATE_NAMES apocytochrome b
ORGANISM #formal_name Allomyces macrogynus
DATE 28-Oct-1996 #sequence_revision 27-Feb-1997 #text_change 08-Sep-1997
ACCESSIONS S63638
REFERENCE S63635

#authors Paguin, B.; Lang, B.F.
#journal J. Mol. Biol. (1996) 255:688-701
#title The mitochondrial DNA of *Allomyces macrogynus*: the complete
#cross-references EMBL:U41288; NID:g1236403; PID:g1236404
#accession S63638
#status nucleic acid sequence not shown; translation not shown
#molecule-type DNA
#residues 1-382 ##label PAQ
#cross-references EMBL:U41288; NID:g1236403; PID:g1236404
#note The nucleotide sequence was submitted to the EMBL Data
Library, November 1995

GENETICS
#gene cob
#introns mitochondrion
#introns 67/3; 137/3; 143/3; 164/1; 200/2; 252/3
#superfamily cytochrome b; cytochrome b homology; cytochrome
CLASSIFICATION b6 homology; plastocyanin--plastocyanin reductase 17K
protein homology
mitochondrion; oxidoreductase

KEYWORDS
FEATURE 10-340 #domain cytochrome b homology #label CBH\
10-210 #domain cytochrome b6 homology #label CB6\
222-340 #domain plastocyanin--plastocyanin reductase 17K protein
homology #label 17K

SUMMARY
#length 382 #molecular-weight 43467 #checksum 6973

Query Match 16.7%; Score 93; DB 2; Length 382;
Best Local Similarity 33.3%; Pred. No. 8.03e-01;
Matches 17; Conservative 15; Mismatches 16; Indels 3; Gaps 3;

Db 8 PVLSTLNSFLDPSPLNITVLMNFGSLG-LCLVIOIVYAGVLAHMYAP 57
1-23
22 PGMITDLK-LNEFTL-TNSYESFLLEMLKLLHLHSGSVTLHHRAS 70

RESULT 3
ENTRY A48156 #type complete
TITLE translation regulator GCD6 - yeast (*Saccharomyces cerevisiae*)
ALTERNATE_NAMES guanine nucleotide exchange factor chain GCD6; protein
YD8142B.03; protein YD8114v; translation initiation factor
eIF-2B homolog

ORGANISM #formal_name *Saccharomyces cerevisiae*
DATE 28-May-1993 #sequence_revision 03-May-1994 #text_change
05-Feb-1998

ACCESSIONS
REFERENCE A48156; S61578; S30776
#authors Bushman, J.L.; Asuru, A.I.; Mats, R.L.; Hinnebusch, A.G.
#journal Mol. Cell. Biol. (1993) 13:1920-1932
#title Evidence that GCD6 and GCD7, translational regulators of
GCN4, are subunits of the guanine nucleotide exchange
factor for eIF-2 in *Saccharomyces cerevisiae*.
#cross-references MUID:93180841
#accession A48156
#molecule-type DNA
#residues 1-712 ##label BUS
#cross-references EMBL:L07115; NID:g11572; PID:g11574
#note sequence extracted from NCBI backbone (NCBIN:126018,
NCBIP:126021)

REFERENCE S61576
#authors Oliver, K.; Harris, D.
#submision submitted to the EMBL Data Library, December 1995
#accession S61578
#molecule-type DNA
#residues 1-712 ##label OLI
#cross-references EMBL:Z68195; NID:g1122341; PID:e213795; PID:g1122344;
MIPS:YDR211v

GENETICS
#gene SCD:GCD6
#cross-references SCD:S0002619; MIPS:YDR211w
#map_position 4R
KEYWORDS translation regulation

SUMMARY #length 712 #molecular-weight 81160 #checksum 142

Query Match 16.5%; Score 92; DB 2; Length 712;
Best Local Similarity 39.5%; Pred. No. 1.07e+00;
Matches 17; Conservative 8; Mismatches 14; Indels 4; Gaps 4;

Db 12 LGN-HGKNSDMPEDRLQAVYLDTS-YETREMFPLAVKPRCLL 52
1-23
14 LGSSHGTCGEMTLQ-LKLESFLTNSSYESSEFLLEL-KL-CLL 54

RESULT 4
ENTRY JC5237 #type complete
TITLE osmotin-like protein precursor - tomato
ORGANISM #formal_name *Lycopersicon esculentum* #common_name tomato
DATE 13-Mar-1997 #sequence_revision 13-Mar-1997 #text_change
13-Nov-1998

ACCESSIONS
REFERENCE JC5237
#authors Chen, R.; Wang, F.; Smith, A.G.
#journal Gene (1996) 179:301-302
#title A flower-specific gene encoding an osmotin-like protein from
Lycopersicon esculentum.
#cross-references MUID:97128324
#accession JC5237
#contents flower
#status preliminary; nucleic acid sequence not shown

FEATURE 1-23
1-23 #domain signal sequence #status predicted #label sig\
24-252 #product osmotin-like protein #status predicted #label
MAT

SUMMARY #length 252 #molecular-weight 27265 #checksum 2939

Query Match 16.3%; Score 91; DB 2; Length 252;
Best Local Similarity 54.2%; Pred. No. 1.41e+00;
Matches 13; Conservative 4; Mismatches 7; Indels 0; Gaps 0;

Db 10 LPLSLFTLTLSSQSTNPFLTL 33
1-11
5 LPLVLLTLTLGSSHGTCGEMTLQ 28

RESULT 5
ENTRY S73757 #type complete
TITLE hypothetical protein FLI_0r1879 - *Mycoplasma pneumoniae* (ATCC
29342) (SGC3)
ORGANISM #formal_name *Mycoplasma pneumoniae*
DATE 27-Feb-1997 #sequence_revision 25-Apr-1997 #text_change
17-Jul-1998

ACCESSIONS
REFERENCE S73757
#authors Himmerle, R.; Hilbert, H.; Plagens, H.; Pirkl, E.; Li,
B.C.; Herrmann, R.
#journal Nucleic Acids Res. (1996) 24:4420-4449
#title Complete sequence analysis of the genome of the bacterium
Mycoplasma pneumoniae.
#cross-references MUID:97105885
#accession S73757
#status preliminary; nucleic acid sequence not shown;
translation not shown

GENETICS
#molecule-type DNA
#residues 1-879 ##label HIM
#cross-references EMBL:AE000042; GB:U00089; NID:g1674112; PID:g1674117
#note the nucleotide sequence was submitted to the EMBL Data
Library, November 1996

GENETICS
#genetic_code SGC3

[illegible]

DE HYPOTHETICAL HELICASE K12H4.8 IN CHROMOSOME III.
 GN K12H4.8.
 OS CAENORABDITIS ELEGANS.
 OC EUKARYOTA; METAZOA; NEMATODA; SECERNENTERA; RHABDITIA; RHABDITIDA;
 CC RHABDITINA; RHABDITOIDEA; RHABDITIDAE; PELODERINAE; CAENORABDITIS.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BRISTOL N2;
 RX MEDLINE: 94150718.
 RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BEKKS M.,
 RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
 RA CRATON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FRASER A.,
 RA FULTON L., GARDNER A., GREEN P., HAMKINS T., HILLIER L., JIER M.,
 RA JOHNSTON L., JONES M., KERSHAW J., KIRSTEN J., LAISTER N.,
 RA LATREILLE P., LIGHTNING J., LLOYD C., MORTIMORE B., O'CALLAGHAN M.,
 RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
 RA SING M., SMALDON N., SMITH A., SMITH M., SONNHAMMER E., STADEN R.,
 RA SULSTON J., THIERRY-MIEG J., THOMAS K., VANDIN M., VAUGHAN K.,
 RA WATSON R., WATSON A., WEINSTOCK L., WILKINSON-SPROAT J.,
 RA WOHLDMAN P.,
 RT "2.2 kb of contiguous nucleotide sequence from chromosome III of C.
 RT elegans.";
 RL NATURE 368:32-38(1994).
 CC -1- SIMILARITY: WITH OTHER ATP DEPENDENT HELICASES.
 CC -1- SIMILARITY: CONTAINS A RNASE III DOMAIN.
 CC
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 CC
 DR EMBL: L14331; G289703; -
 DR PIR: S44849; S44849
 DR WORMPEP: K12H4.8; CE00273.
 DR PROSITE: PS00517; RIBONUCLEASE_III, 1.
 DR PFAM: PF00035; dsrm, 1.
 DR PFAM: PF00271; helicase_C, 1.
 DR PFAM: PF00636; Ribonuclease_3, 2.
 DR KW HYPOTHETICAL PROTEIN; HELICASE; ATP-BINDING; HYDROLASE; NUCLEASE;
 KW ENDONUCLEASE.
 FT NP_BIND 33 40 ATP (POTENTIAL).
 FT SITE 145 148 DECH BOX.
 FT DOMAIN 1554 1822 RNASE III DOMAIN.
 FT SEQUENCE 1822 AA; 208291 MW; 4F856BE CRC32;
 SQ
 Query Match 15.8%; Score 88; DB 1; Length 1822;
 Best Local Similarity 31.4%; Pred. No. 1.13e+00;
 Matches 16; Conservative 17; Mismatches 15; Indels 3; Gaps 3;
 Db 1300 IGLVSPCLLTALTTLSNAD-GMSLEFETIGDSFLKFAITDTLVHTLLD 1349
 QY 1 WGSGLPLVLLTLSSHGSGPMTQ-L-KIKESFLNLSYESSFLEELLE 49
 RESULT 5
 ID RFLN_SCHPO STANDARD; PRT; 396 AA.
 AC 009691;
 DT 01-NOV-1995 (REL. 32, CREATED)
 DT 01-NOV-1995 (REL. 32, LAST SEQUENCE UPDATE)
 DT 01-NOV-1995 (REL. 32, LAST ANNOTATION UPDATE)
 DE PUTATIVE MITOCHONDRIAL PEPTIDE CHAIN RELEASE FACTOR PRECURSOR.
 GN SPAC257.17.
 OS SCHIZOSACCHAROMYCES POMBE (FISSION YEAST).
 OC EUKARYOTA; FUNGI; ASCOMYCOTA; ARCHASCOMYCETES;
 CC SCHIZOSACCHAROMYCETALES; SCHIZOSACCHAROMYCETACEAE;
 CC SCHIZOSACCHAROMYCES.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-972;
 RA GENTLES S., CHURCHER C.M., BARRELL B.G., RAJANDREAM M.A., WALSH S.V.;

RL SUBMITTED (JUL-1995) TO EMBL/GENBANK/DBJ DATA BANKS.
 CC -1- SIMILARITY: BELONGS TO THE PROKARYOTIC AND MITOCHONDRIAL RELEASE
 CC FACTORS FAMILY. STRONG, TO YEAST MRP-1.
 CC
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 CC
 DR EMBL: Z50142; G1052800; -
 DR PROSITE: PS00745; RE_PROK_I, 1.
 DR PFAM: PF00472; RP-1, 1.
 DR KW HYPOTHETICAL PROTEIN; PROTEIN BIOSYNTHESIS; MITOCHONDRION;
 KM TRANSIT PEPTIDE.
 FT TRANSIT 1 ?
 FT CHAIN ? 396 MITOCHONDRION (POTENTIAL).
 FT POTATIVE MITOCHONDRIAL PEPTIDE CHAIN
 FT RELEASE FACTOR.
 SQ SEQUENCE 396 AA; 44954 MW; 988689CB CRC32;
 Query Match 15.6%; Score 87; DB 1; Length 396;
 Best Local Similarity 45.0%; Pred. No. 1.52e+00;
 Matches 9; Conservative 8; Mismatches 3; Indels 0; Gaps 0;
 Db 281 LTHPIGIVTSMQDSRSQHQ 300
 QY 54 LHLFSGTSVTLHRSQHH 73
 RESULT 6
 ID TYTD_PACSU STANDARD; PRT; 272 AA.
 AC P39063;
 DT 01-FEB-1995 (REL. 31, CREATED)
 DT 01-FEB-1995 (REL. 31, LAST SEQUENCE UPDATE)
 DT 15-DEC-1998 (REL. 37, LAST ANNOTATION UPDATE)
 DE HYPOTHETICAL 30.1 KD PROTEIN IN ACUC 5' REGION (ORFA).
 GN TYTD.
 OS BACILLUS SUBTILIS.
 OC BACTERIA; FIRMICUTES; BACILLUS/CLOSTRIDIUM GROUP; BACILLACEAE;
 CC BACILLUS.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-168;
 RX MEDLINE: 95020526.
 RA GRUNDY F.J., WATERS D.A., TAKOVA T.Y., HENKIN T.M.;
 RT "Identification of genes involved in utilization of acetate and
 RT acetoin in Bacillus subtilis".
 RL MOL. MICROBIOL. 10:259-271(1993).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE: 98048467.
 RA LAPIDUS A., GALLERON N., SOROKIN A., EHRLICH S.D.;
 RT "Sequencing and functional annotation of the Bacillus subtilis genes
 RT in the 200 kb trnB-dnaB region".
 RL MICROBIOLOGY 143:3431-3441(1997).
 CC -1- FUNCTION: MAY BE INVOLVED IN SOME TRANSPORT FUNCTION.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE MOTA FAMILY.
 CC
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 CC
 DR EMBL: L17309; G348048; -
 DR EMBL: AF008220; G2293222; -
 DR EMBL: Z99119; E1185846; -
 DR PIR: S39641; S39641.

RT "Metal-tetracycline/H+ antiporter of Escherichia coli encoded by a
 RT transposon Tn10. Histidine 257 plays an essential role in H+
 RT translocation."
 RT J. BIOL. CHEM. 266:6045-6051(1991).
 RN [4]
 RP MUTAGENESIS OF 65-66.
 RX MEDLINE; 90368753.
 RA YAMAGUCHI A., ONO N., AKASAKA T., NOGMI T., SAMAI T.;
 RT "Metal-tetracycline/H+ antiporter of Escherichia coli encoded by a
 RT transposon, Tn10. The role of the conserved dipeptide, ser65-Asp66,
 RT in tetracycline transport."
 RT J. BIOL. CHEM. 265:15525-15530(1990).
 CC -1- FUNCTION: RESISTANCE TO TETRACYCLINE BY AN ACTIVE TETRACYCLINE
 CC EFFLUX THIS IS AN ENERGY-DEPENDENT PROCESS THAT DECREASES THE
 CC ACCUMULATION OF THE ANTIBIOTIC IN HOST CELLS. THIS PROTEIN
 CC FUNCTIONS AS A METAL-TETRACYCLINE/H+ ANTIPORTER.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE.
 CC -1- SIMILARITY: BELONGS TO THE MAJOR FACILITATOR FAMILY (ALSO KNOWN
 CC AS THE DRUG RESISTANCE TRANSDUCASE FAMILY).
 CC -----
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 CC -----
 DR EMBL; V00611; G43701; -;
 DR EMBL; J01830; G154847; -;
 DR PIR; A03507; YBECT0.
 DR PROSITE; P500216; SUGAR_TRANSPORT_1; UNKNOWN.1.
 DR ANTI-BIOTIC RESISTANCE; TRANSMEMBRANE; INNER MEMBRANE; TRANSPORT;
 KW SYMPTOT; TRANSPOSABLE ELEMENT.
 KM
 FT TRANSSEM 6 26
 FT TRANSSEM 42 62 POTENTIAL.
 FT TRANSSEM 74 94 POTENTIAL.
 FT TRANSSEM 102 122 POTENTIAL.
 FT TRANSSEM 131 151 POTENTIAL.
 FT TRANSSEM 159 179 POTENTIAL.
 FT TRANSSEM 214 234 POTENTIAL.
 FT TRANSSEM 244 265 POTENTIAL.
 FT TRANSSEM 277 297 POTENTIAL.
 FT TRANSSEM 298 318 POTENTIAL.
 FT TRANSSEM 336 356 POTENTIAL.
 FT TRANSSEM 363 383 POTENTIAL.
 FT TRANSSEM 65 65 S->C: ALMOST NO CHANGE IN ACTIVITY.
 FT TRANSSEM 65 65 S->A: NO CHANGE IN ACTIVITY.
 FT TRANSSEM 65 65 D->N: UNABLE TO EXTITUDE TETRACYCLINE.
 FT TRANSSEM 65 65 D->E: MODERATE RESISTANCE TO
 FT TRANSSEM 66 66 TETRACYCLINE.
 FT TRANSSEM 257 257 H->E: NO H+ TRANSLLOCATION.
 FT TRANSSEM 257 257 G->D: E (IN REF. 2).
 FT TRANSSEM 281 281 V -> D (IN REF. 2).
 FT TRANSSEM 301 301 Q -> E (IN REF. 2).
 FT TRANSSEM 330 330 A -> T (IN REF. 2).
 FT TRANSSEM 354 354
 SQ SEQUENCE 401 AA; 43267 MW; 4823C395 CRC32;
 Query Match 14.9%; Score 83; DB 1; Length 401;
 Best Local Similarity 31.0%; Pred. No. 4,90e+00;
 Matches 18; Conservative 13; Mismatches 23; Indels 4; Gaps 4;
 Db 1 MNSSTIAVITLDM-GIGLIMPVPLTLE-FlASEDIANHEVLA-LYALMOV 55
 Oy 1 MGSLPLVLTLLSSHGTPGPT-LQIKESFTNSSESSFLEIKLCLHL 57
 RESULT 10 STANDARD: PRT; 445 AA.
 ID YGCS_ECOLI
 AC 046909;
 DT 01-NOV-1997 (REL. 35, CREATED)
 DT 01-NOV-1997 (REL. 35, LAST SEQUENCE UPDATE)

DT 01-NOV-1997 (REL. 35, LAST ANNOTATION UPDATE)
 DE HYPOTHEICAL METABOLITE TRANSPORT PROTEIN IN CISJ-ENO INTERGENIC
 DE REGION.
 GN YGCS.
 OS ESCHERICHIA COLI.
 OC BACTERIA; PROTEOBACTERIA; GAMMA SUBDIVISION; ENTEROBACTERIACEAE;
 OC ESCHERICHIA.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-K12 / MC1655.
 RX MEDLINE; 97426617.
 RA BLATTNER F.R., PLUNKETT G. III, BLOCH C.A., PERNA N.T., BURLAND V.,
 RA RILEY M., COLLADO-VIDES J., GLASNER F.D., ROBE C.R., MAYHEW G.F.,
 RA GREGOR J., DAVIS N.W., KIRKPATRICK H.A., GOEDEN M.A., ROSE D.J.,
 RA MAU B., SHAO Y.;
 RT "The complete genome sequence of Escherichia coli K-12."
 RT SCIENCE 277:1453-1474(1997).
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE
 CC (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE SUGAR TRANSPORTER FAMILY.
 CC -----
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 CC -----
 DR EMBL; U29579; G882664; ALT_INIT.
 DR EMBL; AEO00360; G1789130; ALT_INIT.
 DR ECOGENE; EG13126; YGCS.
 DR PROSITE; P500216; SUGAR_TRANSPORT_1; FALSE_NEG.
 DR PROSITE; P500217; SUGAR_TRANSPORT_2; 1.
 DR PRAM; PF00083; sugar tr. 1
 DR HYPOTHEICAL PROTEIN; TRANSPORT; TRANSMEMBRANE; INNER MEMBRANE.
 KM
 FT TRANSSEM 23 43 POTENTIAL.
 FT TRANSSEM 57 77 POTENTIAL.
 FT TRANSSEM 86 106 POTENTIAL.
 FT TRANSSEM 115 135 POTENTIAL.
 FT TRANSSEM 143 163 POTENTIAL.
 FT TRANSSEM 176 196 POTENTIAL.
 FT TRANSSEM 254 274 POTENTIAL.
 FT TRANSSEM 287 307 POTENTIAL.
 FT TRANSSEM 312 332 POTENTIAL.
 FT TRANSSEM 338 358 POTENTIAL.
 FT TRANSSEM 370 390 POTENTIAL.
 FT TRANSSEM 401 421 POTENTIAL.
 SQ SEQUENCE 445 AA; 48234 MW; BDD078EF CRC32;
 Query Match 14.9%; Score 83; DB 1; Length 445;
 Best Local Similarity 38.6%; Pred. No. 4,90e+00;
 Matches 17; Conservative 9; Mismatches 16; Indels 2; Gaps 2;
 Db 296 GALLGLVLTLLAHRKFLGSLLAIVVACIPSGSSITL 338
 Oy 23 GMTLQIKIKESFTNSSESSFLEIKLCLHLHPGTVTL 65
 RESULT 11 STANDARD: PRT; 647 AA.
 ID BGLI_MOUSE
 AC P23780;
 DT 01-NOV-1991 (REL. 20, CREATED)
 DT 01-NOV-1991 (REL. 20, LAST SEQUENCE UPDATE)
 DT 01-NOV-1997 (REL. 35, LAST ANNOTATION UPDATE)
 DE BETA-GALACTOSIDASE PRECURSOR (EC 3.2.1.23) (LACTASE) (ACID BETA-
 DE GALACTOSIDASE).
 GN GBL1 OR GBL-1 OR BGL.
 OS MUS MUSCULUS (MOUSE).
 OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; MAMMALIA; EUTHERIA;
 OC RODENTIA; SCIURIGRAPHI; MURIDAE; MURINAE; MUS.
 RN [1]
 RP SEQUENCE FROM N.A.

RC TISSUE-BRAIN;
 RX MEDLINE: 91076843.
 RA NAKABA E., SUZUKI K.;
 RT "Molecular cloning of mouse acid beta-galactosidase cDNA: sequence,
 RT expression of catalytic activity and comparison with the human
 RT enzyme.";
 RL BIOCHEM. BIOPHYS. RES. COMMUN. 173:141-148(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-DBA/2J;
 RX MEDLINE: 91298941.
 RA NAKABA E., SUZUKI K.;
 RT "Organization of the mouse acid beta-galactosidase gene.";
 RL BIOCHEM. BIOPHYS. RES. COMMUN. 178:158-164(1991).
 CC -1- FUNCTION: CLEAVES BETA-LINKED TERMINAL GALACTOSYL RESIDUES FROM
 CC GANGLIOSIDES, GLYCOPROTEINS, AND GLYCOSAMINOGLYCAN.
 CC -1- CATALYTIC ACTIVITY: HYDROLYSIS OF TERMINAL, NON-REDUCING BETA-D-
 CC GALACTOSE RESIDUES IN BETA-D-GALACTOSIDES.
 CC -1- SIMILARITY: BELONGS TO FAMILY 35 OF GLYCOSYL HYDROLASES.
 CC -----
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 CC -----
 DR EMBL: M57734; G192187; -
 DR EMBL: M75122; G192185; -
 DR EMBL: M75137; G192185; JOINED.
 DR EMBL: M75107; G192185; JOINED.
 DR EMBL: M75108; G192185; JOINED.
 DR EMBL: M75109; G192185; JOINED.
 DR EMBL: M75111; G192185; JOINED.
 DR EMBL: M75112; G192185; JOINED.
 DR EMBL: M75113; G192185; JOINED.
 DR EMBL: M75114; G192185; JOINED.
 DR EMBL: M75115; G192185; JOINED.
 DR EMBL: M75116; G192185; JOINED.
 DR EMBL: M75117; G192185; JOINED.
 DR EMBL: M75118; G192185; JOINED.
 DR EMBL: M75119; G192185; JOINED.
 DR EMBL: M75120; G192185; JOINED.
 DR EMBL: M75121; G192185; JOINED.
 DR PIR: A37086; A37086.
 DR MGD: MGT:88151; BGL.
 DR PROSITE: PS01182; GLYCOSYL_HYDROL_F35; 1.
 DR PRAM: PF01301; Glycosyl_Hydrl17; 1.
 KW HYDROLASE; GLYCOSIDASE; LYSOSOME; SIGNAL; GLYCOPROTEIN.
 FT SIGNAL 1 24
 FT PROPEP 25 29
 FT CHAIN 30 647
 FT ACT_SITE 189 189 BETA-GALACTOSIDASE.
 FT ACT_SITE 269 269 PROTON DONOR (POTENTIAL).
 FT CARBOHYD 268 268 NUCLEOPHILE (POTENTIAL).
 FT CARBOHYD 268 268 POTENTIAL.
 FT CARBOHYD 500 500 POTENTIAL.
 FT CARBOHYD 504 504 POTENTIAL.
 FT CARBOHYD 510 510 POTENTIAL.
 FT CARBOHYD 544 544 POTENTIAL.
 FT CARBOHYD 557 557 POTENTIAL.
 FT CARBOHYD 617 617 POTENTIAL.
 FT CONFLICT 517 517 N -> D (IN REF. 2).
 FT CONFLICT 539 539 G -> R (IN REF. 2).
 SO SEQUENCE 647 AA; 73121 MW; 15BCFL38 CRC32;
 Query Match 14.9%; Score 83; DB 1; Length 647;
 Best Local Similarity 37.8%; Pred. No. 4.90e+00;
 Matches 14; Conservative 8; Mismatches 12; Indels 3; Gaps 3;

Qy 5 LPIVLLTLTGSSHG-T-G-PGNTLQKL-RESPTNS 38
 ID NME3_HUMAN STANDARD; PRT; 1233 AA.
 AC 014957;
 DT 01-NOV-1997 (REL. 35, CREATED)
 DT 01-NOV-1997 (REL. 35, LAST SEQUENCE UPDATE)
 DT 01-NOV-1997 (REL. 35, LAST ANNOTATION UPDATE)
 DE GLUTAMATE [NMDA] RECEPTOR SUBUNIT EPSILON 3 PRECURSOR (N-METHYL
 DE D-ASPARTATE RECEPTOR SUBTYPE 2C) (NR2C) (NMDAR2C).
 GN GRIN2C.
 OS HOMO SAPIENS (HUMAN).
 OC EUMETAZOA; METAZOA; CHORDATA; VERTEBRATA; MAMMALIA; EUTHERIA;
 OC PRIMATES; CATARRHINI; HOMINIDAE; HOMO.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA TISSUE-BRAIN;
 RA LIN Y.J., BOVERTO S., CARVER J., GIORDANO T.;
 RL SUBMITTED (FEB-1996) TO EMBL/GENBANK/DBJ DATA BANKS.
 CC -1- FUNCTION: NMDA RECEPTOR SUBTYPE OF GLUTAMATE-GATED ION CHANNELS
 CC POSSESSES HIGH CALCIUM PERMEABILITY AND VOLTAGE-DEPENDENT
 CC SENSITIVITY TO MAGNESIUM AND IS MEDIATED BY GLYCINE.
 CC -1- SUBUNIT: HETERODIMER OF AN EPSILON SUBUNIT AND A ZETA SUBUNIT.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
 CC -1- SIMILARITY: BELONGS TO THE LIGAND-GATED IONIC CHANNELS FAMILY.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: L76224; G1196449; -
 DR MIM: L38254; -
 DR PRAM: PF00060; Lig_chan; 1.
 KW RECEPTOR; SIGNAL; TRANSMEMBRANE; POSTSYNAPTIC MEMBRANE; CALCIUM;
 KW IONIC CHANNEL; MAGNESIUM.
 FT SIGNAL 1 19
 FT CHAIN 20 1233
 FT TRANSMEM 554 574 1 (POTENTIAL).
 FT TRANSMEM 597 617 2 (POTENTIAL).
 FT TRANSMEM 627 647 3 (POTENTIAL).
 FT TRANSMEM 815 835 4 (POTENTIAL).
 FT SITE 612 612 FUNCTIONAL DETERMINANT OF NMDA
 FT CARBOHYD 70 70 RECEPTORS (BY SIMILARITY).
 FT CARBOHYD 337 337 POTENTIAL.
 FT CARBOHYD 438 438 POTENTIAL.
 FT CARBOHYD 539 539 POTENTIAL.
 SO SEQUENCE 1233 AA; 134239 MW; 671F9981 CRC32;
 Query Match 14.9%; Score 83; DB 1; Length 1233;
 Best Local Similarity 33.3%; Pred. No. 4.90e+00;
 Matches 20; Conservative 13; Mismatches 21; Indels 3; Gaps 3;
 Db 1 MGSGPLVLLT-SFGAMGLPGGEGCGMTVAVFSSGCPPOAFARLTPOSFL-DIP 59
 Qy 1 MGSGPLVLLT-LTSSHGTPGNTLQ-LKLRESPTNSYSESFLELLEKLCILHLIP 58
 RESULT 13
 ID INVE_SALTY STANDARD; PRT; 372 AA.
 AC P35671;
 DT 01-JUN-1994 (REL. 29, CREATED)
 DT 01-FEB-1995 (REL. 31, LAST SEQUENCE UPDATE)
 DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)
 DE INVASION PROTEIN INVE.
 GN INVE.

OS SALMONELLA TYPHIMURIUM.
OC BACTERIA: PROTEOBACTERIA: GAMMA SUBDIVISION: ENTEROBACTERIACEAE;
OC SALMONELLA.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-SR11 / SL1344;
RX MEDLINE: 92335220.
RA GINOCCHIO C., PACE J., GALAN J.E.;
RT "Identification and molecular characterization of a Salmonella typhimurium gene involved in triggering the internalization of salmonellae into cultured epithelial cells."
RL PROC. NATL. ACAD. SCI. U.S.A. 89:5976-5980(1992).
RN [2]
RP SEQUENCE OF 1-69 FROM N.A.
RC STRAIN-SR11 / SL1344;
RX MEDLINE: 95089692.
RA KANIGA K., BOSSIO J.C., GALAN J.E.;
RT "The Salmonella typhimurium invasion genes invF and invG encode homologues of the *Yersinia* and *Pseudomonas* family of proteins."
RL MOL. MICROBIOL. 13:555-568(1994).
RN [3]
RP SEQUENCE OF 1-5 FROM N.A.
RC STRAIN-TML;
RA LODGE J.M., AMIN I.I., DOUCE G.R., BROWN N.L., STEPHEN J.;
RT SUBMITTED (SEP-1993) TO EMBL/GENBANK/DBJ DATA BANKS.
CC -1- FUNCTION: INVOLVED IN THE TRIGGERING OF INTRACELLULAR EVENTS THAT LEAD TO MICROBIAL INTERNALIZATION OF THE INTESTINAL EPITHELIUM. THESE EVENTS INCLUDE INCREASE IN CALCIUM LEVEL, REDISTRIBUTION OF ACTIN MICROFILAMENTS, AND CHANGES IN THE NORMAL STRUCTURE OF THE MICROVILLI.
CC -1- SIMILARITY: TO YERSINIA OUTER MEMBRANE PROTEIN YOPN (LCRE).
CC -----
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CC -----
DR EMBL: M90714; -; NOT_ANNOTATED_CDS.
DR EMBL: U08280; G497226; -;
DR EMBL: X75302; E86775; ALT_FRAME.
DR STYGENE: SG10187; INVE.
KW VIRULENCE.
SQ SEQUENCE 372 AA: 42435 MW: A345002E CRC32:

Query Match 14.7%; Score 82; DB 1; Length 372;
Best Local Similarity 26.1%; Pred. No. 6.52e+00;
Matches 12; Conservative 17; Mismatches 16; Indels 1; Gaps 1;

Db 284 LILMSLILQPHVDLSLADIGLNLILSHKEH-ASFLDIFQVOC 328
QY 7 LVLLTLGLSSHGTPGPGMTQLKLSKESFLTNSYSSLSLELLKLC 52

RT niger WT-2223L."
RL SUBMITTED (JUL-1995) TO EMBL/GENBANK/DBJ DATA BANKS.
RN [2]
RP SEQUENCE OF 142-286 FROM N.A.
RC STRAIN-IFM 5367, IFM 5368, IFM 40606, IFM 41398, IFM 41399, IFM 46897;
RX MEDLINE: 99065785.
RA WANG L., YOKOYAMA K., MIYAJI M., NISHIMURA K.;
RT "The identification and phylogenetic relationship of pathogenic species of *Aspergillus* based on the mitochondrial cytochrome b gene."
RL MCOL. 36:153-164(1996).
CC -1- FUNCTION: COMPONENT OF THE UBIQUINOL-CYTOCHROME C REDUCTASE COMPLEX (COMPLEX III OR CYTOCHROME B-C1 COMPLEX), WHICH IS A RESPIRATORY CHAIN THAT GENERATES AN ELECTROCHEMICAL POTENTIAL COUPLED TO ATP SYNTHESIS.
CC -1- CATALYTIC ACTIVITY: OX(2) + 2 FERRICYTOCHROME C
CC -1- CORRECTOR: TWO HEME GROUPS (B562 AND B566) WHICH ARE NOT COVALENTLY BOUND TO THE PROTEIN.
CC -1- SUBUNIT: THE MAIN SUBUNITS OF COMPLEX B-C1 ARE: CYTOCHROME B, CYTOCHROME C1 AND THE RIESKE PROTEIN.
CC -----
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CC -----
DR EMBL: D63375; G1000910; -;
DR EMBL: AB000575; D1035119; -;
DR EMBL: AB000576; D1035120; -;
DR EMBL: AB000577; D1035121; -;
DR EMBL: AB000578; D1035122; -;
DR EMBL: AB000583; D1035127; -;
DR EMBL: AB000597; D1035139; -;
DR PROSITE: PS00192; CYTOCHROME_B_HEME: 1.
DR PROSITE: PS00193; CYTOCHROME_B_OO: 1.
DR PFM: PFM0033; Cytochrome B; 1.
DR PFM: PFM0033; Cytochrome B; 1.
KW ELECTRON TRANSPORT; MITOCHONDRION; RESPIRATORY CHAIN; TRANSMEMBRANE;
KW HEME.
FT METAL 82 82 IRON 1 (HEME B562 AXIAL LIGAND).
FT METAL 96 96 IRON 2 (HEME B566 AXIAL LIGAND).
FT METAL 183 183 IRON 2 (HEME B562 AXIAL LIGAND).
FT METAL 197 197 IRON 1 (HEME B566 AXIAL LIGAND).
SQ SEQUENCE 385 AA: 43078 MW: 17050CE1 CRC32:

Query Match 14.7%; Score 82; DB 1; Length 385;
Best Local Similarity 34.3%; Pred. No. 6.52e+00;
Matches 12; Conservative 11; Mismatches 11; Indels 1; Gaps 1;

Db 24 ANISTLVNFGSLA-LGIGQIVTGVTLAMHYTPS 57
QY 36 TNSYSESELELEKLCILHLPLSGTSTVLHHARS 70

RESULT 14
ID CYB.ASPNG STANDARD: PRT: 385 AA.
AC 033798;
DT 15-DEC-1998 (REL. 37, CREATED)
DT 15-DEC-1998 (REL. 37, LAST SEQUENCE UPDATE)
DE CYTOCHROME B (EC 1.10.2.2).
GN CDB OR CYTB OR COBA.
OS ASPERGILLUS NIGER.
OC MITOCHONDRION.
OC EUKARYOTA: FUNGI: ASCOMYCOTA: EUASCOMYCETES: PLBCTOMYCETES:
OC EURHOTIALES: TRICHOCOMACEAE: MITOSPORIC TRICHOCOMACEAE: ASPERGILLUS.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-WO-2223L;
RA NARUSAMA T., KANAYAMA S., KIRIMURA K., USAMI S.;
RT "Nucleotide sequence of the apocytochrome b gene of *Aspergillus*

RESULT 15
ID MOTA.RHOH STANDARD: PRT: 253 AA.
AC 053174;
DT 15-JUL-1998 (REL. 36, CREATED)
DT 15-JUL-1998 (REL. 36, LAST SEQUENCE UPDATE)
DE 15-DEC-1998 (REL. 37, LAST ANNOTATION UPDATE)
DE CHEMOTAXIS MOTA PROTEIN (MOTILITY PROTEIN A).
GN MOTA.
OS RHODOBACTER SPHAEROIDES (RHODOSPIRIONAS SPHAEROIDES).
OC BACTERIA: PROTEOBACTERIA: ALPHA SUBDIVISION: RHODOBACTER GROUP:
OC RHODOBACTER.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-WS8;
RX MEDLINE: 96123438.
RA SHAH D.S.H., SOCKERT R.E.;

```
RT "Analysis of the motA flagellar motor gene from Rhodospirillum rubrum, a bacterium with a unidirectional, stop-start flagellum".
RT flagellum".
RL MOL. MICROBIOL. 17:961-969(1995).
CC -1- FUNCTION: REQUIRED FOR ROTATION OF THE FLAGELLAR MOTOR. PROBABLE
CC TRANSMEMBRANE PROTON CHANNEL (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE.
CC -1- SIMILARITY: BELONGS TO THE MOTA FAMILY.
CC -----
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC -----
DR EMBL: X85986; G758646; -.
DR PROSITE: PS01307; MOT_A: 1.
KW CHEMOTAXIS; FLAGELLA; TRANSMEMBRANE; INNER MEMBRANE;
KW FLAGELLAR ROTATION; HYDROGEN ION TRANSPORT.
FT TRANSMEM 3 23 POTENTIAL.
FT TRANSMEM 29 49 POTENTIAL.
FT TRANSMEM 146 166 POTENTIAL.
FT TRANSMEM 181 201 POTENTIAL.
FT DOMAIN 202 253 CYTOPLASMIC (POTENTIAL).
SQ SEQUENCE 253 AA; 27194 MW; F594C09C CRC32;

Query Match 14.58; Score 81; DB 1; Length 253;
Best Local Similarity 27.58; Pred. No. 8.66e+00;
Matches 14; Conservative 20; Mismatches 14; Indels 3; Gaps 2;

Db 156 VGTLLGLVLMGNMSPKSGIPAMVALITLYGALMANVIF-APLNKLE 205
Oy 1 MSGGLPLVLLLTLLGSSHGTPMTLQL-KLKSEFLTNSYSSSELELLE 49
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Search completed: Fri Oct 22 18:43:49 1999
Job time : 22 secs.

OY 32 ESFLTNSYSSFFLELKLCLLHLPGSTVTLHAR 69

RESULT 6 PRELIMINARY; PRT: 297 AA.
AC 003305;
DT 01-JUL-1997 (TREMBLREL. 04, CREATED)
DT 01-JUL-1997 (TREMBLREL. 04, LAST SEQUENCE UPDATE)
DT 01-NOV-1996 (TREMBLREL. 01, LAST SEQUENCE UPDATE)
DE CYTOCHROME B (FRAGMENT)
OS CAROTHOCHLYS INSCULPIA (PITTED-SHELLED TURTLE).
OC MITOCHONDRION
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TESTUDINES; CRYPTODIRA;
OC TRIONCHOIDEA; CAROTHOCHLYDAE; CAROTHOCHLYS.
RN [1]
RP SEQUENCE FROM N.A.
NA SHAFFER H.B., MEYLAN P., MCKNIGHT M.L.;
RL SYST. BIOL. 0:0-0(0).
CC -1- CATALYTIC ACTIVITY: OH(2) + 2 FERRICYTOCHROME C = Q + 2
CC -1- FERROCYTOCHROME C
CC -1- COFACTOR: TWO HEME GROUPS
CC (B562 AND B566) WHICH ARE NOT COVALENTLY BOUND TO THE PROTEIN
CC (BY SIMILARITY)
DR EMBL: U01355; G2098653;
DR PROSITE: PS00192; CYTOCHROME-B-HEME; 1.
DR PFAM: PF00032; cytochrome-b_c; 1.
DR PFAM: PF00033; cytochrome-b_n; 1.
KM MITOCHONDRION; ELECTRON TRANSPORT; RESPIRATORY CHAIN; TRANSMEMBRANE;
KW HEME.
FT NON_TER 1 1
FT NON_TER 297 297
SQ SEQUENCE 297 AA; 33587 MW; 7D5808B5 CRC32;

Query Match 15.8%; Score 88; DB 8; Length 297;
Best Local Similarity 31.5%; Pred. No. 3.63e+00;
Matches 17; Conservative 10; Mismatches 25; Indels 2; Gaps 2;
DB 165 GLIIVLLFLYETGSNNPTGNSNM-D-KIPFPIYSIKDF-GLIMLAAILNL 216
OY 4 GLPLVLLTLGSSHGTPGNTLQKLESFLTNSYSSFFLELKLCLLHL 57

RESULT 7 PRELIMINARY; PRT: 143 AA.
AC 069582;
DT 01-NOV-1996 (TREMBLREL. 01, CREATED)
DT 01-NOV-1996 (TREMBLREL. 01, LAST SEQUENCE UPDATE)
DT 01-NOV-1996 (TREMBLREL. 08, LAST SEQUENCE UPDATE)
DE HERPESVIRUS TYPE 6 DNA.
OS HUMAN HERPESVIRUS-6.
OC VIRUSES; DSDNA VIRUSES; NO RNA STAGE; HERPESVIRIDAE; BETAHERPESVIRINAE;
OC ROSELOVIRUS.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 94181269.
RA THOMSON J., CHODHURY S., KASHANCHI F., DONIGER J., BERNEMAN Z.,
RA FRENEL N., ROSENTHAL L.J.;
RT "A transforming fragment within the direct repeat region of human
RT herpesvirus type 6 that transactivates HIV-1.";
RL ONCOGENE 9:1167-1175(1994).
DR EMBL: X73675; G469957;
SQ SEQUENCE 143 AA; 13317 MW; 597857A6 CRC32;

Query Match 15.6%; Score 87; DB 14; Length 143;
Best Local Similarity 47.2%; Pred. No. 4.79e+00;
Matches 17; Conservative 4; Mismatches 14; Indels 1; Gaps 1;

DB 88 LGGLGIGLGLGIGLGLGIGLGLGIGLGLAGGFL 123
OY 1 MGSGPLVLLTL-LGSSHGTPGNTLQKLESFL 35

RESULT 8

ID 014968 PRELIMINARY; PRT: 322 AA.
AC 014968;
DT 01-NOV-1996 (TREMBLREL. 01, CREATED)
DT 01-NOV-1996 (TREMBLREL. 01, LAST SEQUENCE UPDATE)
DT 01-NOV-1996 (TREMBLREL. 08, LAST SEQUENCE UPDATE)
DE RHODOPSIN.
OS HOMO SAPIENS (HUMAN).
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; MAMMALIA; EUTHERIA; PRIMATES;
OC CARPARRHINI; HOMINIDAE; HOMO.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 92250505.
RA KUNZ D., GERARD N.P., GERARD C.;
RT "The human leukocyte platelet-activating factor receptor. cDNA
RT cloning, cell surface expression, and construction of a novel
RT epsilon-bearing analog";
RL J. BIOL. CHEM. 267:9101-9106(1992).
DR EMBL: M76676; G189270;
DR PFAM: PF00001; 7tm_1; 2.
KW GTP-BINDING.
SQ SEQUENCE 322 AA; 33096 MW; 70F54EC8 CRC32;

Query Match 15.4%; Score 86; DB 4; Length 322;
Best Local Similarity 34.5%; Pred. No. 6.32e+00;
Matches 19; Conservative 13; Mismatches 22; Indels 1; Gaps 1;
DB 95 ALVLLIIFLSSLNCNMGVYKRRQRTVNAFLIS-LSLSDLITALLCLPAA 148
OY 6 PLVLLTLGSSHGTPGNTLQKLESFLTNSYSSFFLELKLCLLHLPSG 60

RESULT 9 PRELIMINARY; PRT: 562 AA.
AC 023220;
DT 01-NOV-1996 (TREMBLREL. 01, CREATED)
DT 01-NOV-1996 (TREMBLREL. 01, LAST SEQUENCE UPDATE)
DT 01-JAN-1999 (TREMBLREL. 09, LAST SEQUENCE UPDATE)
DE W08D2.3 PROTEIN.
GN W08D2.3.
OS CAENORHABDITIS ELEGANS.
OC EUKARYOTA; METAZOA; NEMATODA; SECERNENTEA; RHABDITIA; RHABDITIDA;
OC RHABDITINA; RHABDITOIDEA; RHABDITIDAE; PELODERINAE; CAENORHABDITIS.
RN [1]
RP SEQUENCE FROM N.A.
RA SWINBURNE J., AINSCOUGH R.;
RL SUBMITTED (MAR-1996) TO EMBL/GENBANK/DBJ DATA BANKS.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE: 94150718.
RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BEKS M.,
RA BONFIELD J., BURTON J., CONNELL M., CORSEY T., COOPER J., COULSON A.,
RA CAXTON M., DEAR S., DU Z., DUBBIN R., FAVELLO A., FULTON L.,
RA GARDNER A., GREEN P., HAWKINS T., HILLER L., JIER M., JOHNSON L.,
RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
RA LIGHTNIE J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
RA SPALDON N., SMITH A., SONNHAMMER E., STADEN R., SLESTON J.,
RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
RA WATSON A., WEINSTOCK L., WILKINSON-SPOAT J., WOHLDMAN P.,
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans";
RL NATURE 366:32-38(1994).
DR EMBL: Z70271; E135011;
SQ SEQUENCE 562 AA; 63331 MW; 5272E400 CRC32;

Query Match 15.4%; Score 86; DB 5; Length 562;
Best Local Similarity 31.5%; Pred. No. 6.32e+00;
Matches 23; Conservative 19; Mismatches 27; Indels 4; Gaps 4;

DB 122 ILLCLVIFVGTGIGLSAIVSN-FVWIF-QQSFL-LIQFLHIGAFSGVAVWFY 178
OY 7 LVLLTLGSSHGTPGNTLQKLESFLTNSYSSFFLELKLCLLHLPGSTVTLH 66

Db 179 QSAFSEHOEDCNS 191
: | | | :
Qy 67 H-ARSOHVCNT 78

RESULT 10
ID 003294: PRELIMINARY: PRT: 297 AA.

DT 01-JUL-1997 (TREMBLREL. 04, CREATED)
DT 01-JUL-1997 (TREMBLREL. 04, LAST SEQUENCE UPDATE)
DT 01-NOV-1998 (TREMBLREL. 08, LAST ANNOTATION UPDATE)
DE CYTOCHROME B (FRAGMENT).
OS HEOSEMYS SPINOSA.
OC MITOCHONDRION.
CC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TESTUDINES; CRYPTODIRA;
CC TESTUDINOIDEA; BATAGURIDAE; HEOSEMYS.
RN [1]
RP SEQUENCE FROM N.A.
RA SHAFFER H.B., MEYLAN P., MCKNIGHT M.L.:
RL SYST. BIOL. 0:0-0(0).
CC -1- CATALYTIC ACTIVITY: QH(2) + 2 FERRICYTOCHROME C = Q + 2
CC FERRICYTOCHROME C.
CC -1- COFACTOR: TWO HEME GROUPS
CC (B562 AND B566) WHICH ARE NOT COVALENTLY BOUND TO THE PROTEIN
(BY SIMILARITY).
DR EMBL: U01362; G2098675; -
DR PROSITE: P800192; CYTOCHROME_B_HEME; 1.
DR PFAM: PF000032; cytochrome_b_c; 1.
DR PFAM: PF000033; cytochrome_b_n; 1.
KW MITOCHONDRION; ELECTRON TRANSPORT; RESPIRATORY CHAIN; TRANSMEMBRANE;
HM HEME.
FT NON_TER 1 1
FT 297 297
SQ SEQUENCE 297 AA; 33321 MW; AD3A6749 CRC32;

Query Match 15.2%; Score 85; DB 8; Length 297;
Best Local Similarity 32.1%; Pred. No. 8.31e+00;
Matches 18; Conservative 11; Mismatches 25; Indels 2; Gaps 2;
Db 164 AGAIYVLLFLHETGSNNPGLNSND-KIPFHPFYKD-LIGLILMLTLTL 217
: | | | | | : | | | | | : | | | | | : | | | | | :
Qy 3 SGLPVLTLTLSSHGTCGMLQKLKESFLNLSYESSFLLEKLCILLHL 58

RESULT 11
ID 003295: PRELIMINARY: PRT: 297 AA.
AC 003295;
DT 01-JUL-1997 (TREMBLREL. 04, CREATED)
DT 01-JUL-1997 (TREMBLREL. 04, LAST SEQUENCE UPDATE)
DT 01-NOV-1998 (TREMBLREL. 08, LAST ANNOTATION UPDATE)
DE CYTOCHROME B (FRAGMENT).
OS TRACHEMYS SCRIPTA (RED-EARED SLIDER TURTLE).
OC MITOCHONDRION.
CC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TESTUDINES; CRYPTODIRA;
CC TESTUDINOIDEA; EMYDIDAE; TRACHEMYS.
RN [1]
RP SEQUENCE FROM N.A.
RA SHAFFER H.B., MEYLAN P., MCKNIGHT M.L.:
RL SYST. BIOL. 0:0-0(0).
CC -1- CATALYTIC ACTIVITY: QH(2) + 2 FERRICYTOCHROME C = Q + 2
CC FERRICYTOCHROME C.
CC -1- COFACTOR: TWO HEME GROUPS
CC (B562 AND B566) WHICH ARE NOT COVALENTLY BOUND TO THE PROTEIN
(BY SIMILARITY).
DR EMBL: U01351; G2098693; -
DR PROSITE: P800192; CYTOCHROME_B_HEME; 1.
DR PFAM: PF000032; cytochrome_b_c; 1.
DR PFAM: PF000033; cytochrome_b_n; 1.
KW MITOCHONDRION; ELECTRON TRANSPORT; RESPIRATORY CHAIN; TRANSMEMBRANE;
HM HEME.
FT NON_TER 1 1
FT 297 297
SQ SEQUENCE 297 AA; 33479 MW; 1BC0CFAC CRC32;

Query Match 15.2%; Score 85; DB 8; Length 297;
Best Local Similarity 33.3%; Pred. No. 8.31e+00;
Matches 18; Conservative 9; Mismatches 25; Indels 2; Gaps 2;

Db 165 GILTVHLLFLHETGSNNPGLNSND-KIPFHPFYKD-LIGLILMLTLTL 216
: | | | | | : | | | | | : | | | | | : | | | | | :
Qy 4 GLPVLTLTLSSHGTCGMLQKLKESFLNLSYESSFLLEKLCILLHL 57

RESULT 12
ID 023674: PRELIMINARY: PRT: 395 AA.
AC 023674;
DT 01-JAN-1998 (TREMBLREL. 05, CREATED)
DT 01-JAN-1998 (TREMBLREL. 05, LAST SEQUENCE UPDATE)
DT 01-NOV-1998 (TREMBLREL. 08, LAST ANNOTATION UPDATE)
DE CHALCONE SYNTHASE HOMOLOG.
GN T7123.4.
OS ARABIDOPSIS THALIANA (MOUSE-EAR CRESS).
OC EUKARYOTA; VIRIDIPANTAE; STEPHOPHYTA; EMERYOPHYTA; TRACHEOPHYTA;
OC EUPHYLLIPHYTES; SPERMATOPHYTA; MAGNOLIOPHYTA; EUDICOTYLEDONS; ROSIDAE;
OC CAPPARALES; BRASSICACEAE; ARABIDOPSIS.
RN [1]
RP SEQUENCE FROM N.A.
RA PEDERSIEL N.A., CONWAY A.B., CONWAY A.R., DAVIS K., BRENDLE V.,
RA PALM C.J., AU M., ARNOLD R., CHUNG E., KURTZ D.B., BUEHLER E.,
RA DEMAR K., FENG J., KIM C., LI Y., SHIN P., SUN H., OUI O.,
RA OSBORNE B., SHEN Y.K., TORIUMI M., VYOTSKAYA V., YU G., THEOLOGIS A.,
RA ECKER J., DAVIS R.W.;
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DR EMBL: U89959; G2317904; -
DR PFAM: PF00195; Chal.stil synt; 1.
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Matches 14; Conservative 12; Mismatches 15; Indels 1; Gaps 1;

Db 27 TLALGKAPSOVPOENLVESFLRDKCDPAFIKKEKLEHC 68
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Qy 12 TLGSSHGTCGMLQKLKESFLNLSYESSFL-ELLEKLC 52

RESULT 13
ID 042453: PRELIMINARY: PRT: 448 AA.
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DT 01-JAN-1998 (TREMBLREL. 05, LAST SEQUENCE UPDATE)
DT 01-NOV-1998 (TREMBLREL. 08, LAST ANNOTATION UPDATE)
DE SERPIN PRECURSOR.
OS PETROMYZON MARINUS (SEA LAMPREY).
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; CEPHALASPIDOMORPHI;
OC PETROMYZONTIFORMES; PETROMYZONTIDAE; PETROMYZON.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-LARVAL LIVER;
RA ROBSON P., YOUSON J.H., KEELLY F.W.;
RL SUBMITTED (JUN-1997) TO EMBL/GENBANK/DBJ DATA BANKS.
DR EMBL: AF009964; G2275601; -
DR PFAM: PF000079; serpin; 1.
KW SIGNBL.
FT SIGNBL 1 21
FT CHAIN 22 448
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SQ SEQUENCE 448 AA; 49044 MW; 6E233618 CRC32;

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RESULT 14
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DT 01-JUN-1998 (TREMBLREL. 06, CREATED)
DT 01-JUN-1998 (TREMBLREL. 06, LAST SEQUENCE UPDATE)
DT 01-NOV-1998 (TREMBLREL. 08, LAST ANNOTATION UPDATE)
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OS CAENORHADITIS ELEGANS.
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RN [1]
RP SEQUENCE FROM N.A.
RA MATTHEWS L.;
RL SUBMITTED (NOV-1996) TO EMBL/GENBANK/DBJ DATA BANKS.
RN [2]
RP SEQUENCE FROM N.A.

RX MEDLINE: 94150718.
RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BEERS M.,
RA BONFIELD J., BURTON J., CONNELL M., COPEY T., COOPER J., COULSON A.,
RA CRAXTON M., DEAR S., DI 2., DORBIN R., FAVELLO A., FULTON L.,
RA GARDNER A., GREEN P., HAKINS T., HILLER L., JER M., JOHNSTON L.,
RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
RA PARSONS J., PERCY C., RIEKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
RA SPALDON N., SMITH A., SONNHAMMER E., STADEN R., SUTTON J.,
RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOLDMAN P.,
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans.";
RL NATURE 368:32-38(1994).
DR EMBL: Z81562; E1188135;
SQ SEQUENCE 676 AA: 77673 MW: 366C5777 CRC32;

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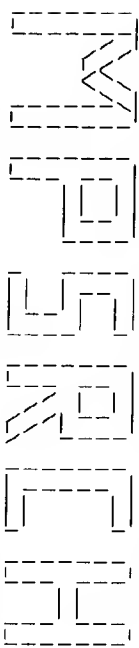
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OY 28 LKLESLFTNNS-YESSFLEIKELICLILHLPSTVTLHH 67

RESULT 15
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AC 043190:
DT 01-JUN-1998 (TREMBLREL. 06, CREATED)
DT 01-JUN-1998 (TREMBLREL. 06, LAST SEQUENCE UPDATE)
DT 01-JUN-1998 (TREMBLREL. 06, LAST ANNOTATION UPDATE)
DE PURINERGIC P2Y11 RECEPTOR.
GN P2Y11.
OS HOMO SAPIENS (HUMAN).
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; MAMMALIA; EUTHERIA; PRIMATES;
OC CATARRHINI; HOMINIDAE; HOMO.
RN [1]
RP SEQUENCE FROM N.A.
RA TISSUE-PLACENTA;
RA COMINI D., GOVAERTS C., PARMENTIER M., BOYNAENS J.M.;
RA J. BIOL. CHEM. 0:0-0(1997).
DR EMBL: AF030335; G2674120;
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OY 1 MGSGPLVLLTLIGS-SHGT--GPGMTQLKKESTLNS 39

Search completed: Fri Oct 22 18:44:57 1999
Job time : 51 secs.



(TM)

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Mparch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 16:51:21 1999; MasPar time 456.53 seconds

Tabular output not generated. 1390.417 Million cell updates/sec

Title: >US-09-092-296-2
Description: (1-229) from US09092296.seq
Perfect Score: 229
N.A. Sequence: 1 ACCGGAGACTGATGTCCTCC.....CCATCTCCCTCAGAGACCA 229
Comp: TGGCCCTGAGAGTCACAGAG.....GGTAGAGGAGAGTCCCTGGT

Scoring table:
Gap 6
TABLE default

Mmatch STD : Dbase 0; Query 0

Searched: 646147 seqs, 1385935633 bases x 2

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database:

emb158
1:em_ba1 2:em_ba2 3:em_fun 4:em_htg 5:em_hum1 6:em_hum2
7:em_in 8:em_com 9:em_or 10:em_ov 11:em_pat 12:em_ph
13:em_pl 14:em_ro 15:em_sts 16:em_vl
genbank111

Database:
17:gb_ba1 18:gb_ba2 19:gb_htg1 20:gb_htg2 21:gb_in1
22:gb_in2 23:gb_com 24:gb_ov 25:gb_pat 26:gb_ph 27:gb_pl1
28:gb_pl2 29:gb_pl3 30:gb_pl4 31:gb_pl5 32:gb_ro
33:gb_stc 34:gb_sts 35:gb_sy 36:gb_un 37:gb_vl

Statistics: Mean 9.569; Variance 4.802; scale 1.993

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description	Pred. No.
1	108	47.2	47323	31	AC0005937	Homo sapiens clone UMG 8.69e-62
2	46	20.1	7218	25	166494	Sequence 14 from patent 9.04e-15
3	33	14.4	215	25	128278	Sequence 5 from patent 4.40e-06
4	33	14.4	216021	31	HUAC004787	Homo sapiens Chromosome 1.30e-03
5	31	13.5	965	25	AR024229	Sequence 22 from patent 7.82e-05
6	29	12.7	74371	31	AC005369	Homo sapiens Chromosome 1.30e-03
7	29	12.7	216021	31	HUAC004787	Homo sapiens Chromosome 1.30e-03
8	27	11.8	215	25	128278	Sequence 5 from patent 1.98e-02
9	26	11.4	1056	23	AR024229	Sequence 22 from patent 7.48e-02
10	26	11.4	1056	23	MY087256	Mustela vison GT dinuc 7.48e-02
11	25	10.9	565	25	E04076	gDNA encoding envelope 2.76e-01
12	25	10.9	6096	31	AC003030	Homo sapiens chromosome 2.76e-01
13	24	10.5	60	25	A62989	Sequence 1 from patent 9.88e-01

C	14	24	10.5	1056	23	MY087256	Mustela vison GT dinuc	9.88e-01
C	15	23	10.0	3290	29	HUMPFUCAS	H.sapiens ficoidase P	3.44e+00
C	16	23	10.0	133457	30	AC003999	Human PAC clone DJ1139	3.44e+00
C	17	23	10.0	203418	19	AC004947	Homo sapiens clone DJ1	3.44e+00
C	18	22	9.6	30	25	A62994	Sequence 6 from patent	1.16e+01
C	19	22	9.6	108	21	D87227	Trypanosoma cruzi mRNA	1.16e+01
C	20	22	9.6	288	29	HS15665R	H.sapiens CPG island D	1.16e+01
C	21	22	9.6	1738	29	HUMGOLGINB	Human (clone SY2/10) 9	1.16e+01
C	22	22	9.6	2180	21	DROSYT	D.melanogaster synapto	1.16e+01
C	23	22	9.6	2273	32	MMBAKEXN	Mus musculus Bak gene,	1.16e+01
C	24	22	9.6	3088	11	E10775	DNA encoding part of A	1.16e+01
C	25	22	9.6	3890	30	AR014574	Sequence 1 from patent	1.16e+01
C	26	22	9.6	4472	17	NSMRBROF	Neisseria meningitidis	1.16e+01
C	27	22	9.6	6640	29	D63987	Homo sapiens mRNA for	1.16e+01
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C	29	22	9.6	79482	19	AP000029	Homo sapiens genomic D	1.16e+01
C	30	22	9.6	82098	31	AC006252	Homo sapiens 3p21.1 co	1.16e+01
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C	37	22	9.6	192853	19	AC005096	Homo sapiens clone RG3	1.16e+01
C	38	22	9.6	241407	20	AC003059	Mouse Chromosome 10 BA	1.16e+01
C	39	22	9.2	66	25	141364	Sequence 143 from pate	3.79e+01
C	40	21	9.2	2976	21	PFAPR2B	Plasmodium knowlesi P-	3.79e+01
C	41	21	9.2	3043	32	AB010141S4	Mus musculus gene for	3.79e+01
C	42	21	9.2	130560	19	HS508115	Human DNA sequence ***	3.79e+01
C	43	21	9.2	149298	19	AP000031	Homo sapiens genomic D	3.79e+01
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TITLE	1	AC005937.1	GI:3845393				
JOURNAL	1						

1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the
GenBank flat file format but are available as part
of this entry's ASN.1 file.

Double stranded (DS) coverage: 75.5%
DS or two chemistry coverage: 98.9%
Single stranded regions: 3

Sequence Validation:

This sequence has been validated by Multiple Complete Digest
Mapping. Comparison of the experimentally derived map digest
fragments with sequence-predicted fragments is given below.
Small fragments below a variable cutoff (approximately 400-600bp)
are not mapped and hence do not appear in the table. There are no
significant remaining discrepancies between the experimental and
predicted values. Uniquely ordered fragment groups are separated
by dashed lines.

Map	Seq	Map	Seq	Map	Seq
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20320.67	20855.00	1050.18	1015.00	3279.08	3231.00
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variation insertion of 17bp repeat

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Query Match 47.2%; Score 108; DB 31; Length 47323;
Best Local Similarity 100.0%; Pred. No. 8.69e-62;
Matches 108; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DEFINITION Sequence 14 from Patent US 5670367.
ACCESSION 166494
NID 92724471
VERSION 166494.1 GI:2724471
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 7218)
AUTHORS Dörner, F., Schefflinger, F. and Falkner, F. Gunter.
TITLE Recombinant fowlpox virus
JOURNAL Patent: US 5670367-A 14 23-SEP-1997;
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repeat_region	6586..6956 /rpt_family="L1" 6647..6684 /note="(CA)19"
repeat_region	/rpt_type=tandem /rpt_unit=CA 7113..7375 /rpt_family="Alu" complement(7830..9185) /standard_name="possible repeat"
repeat_region	8358..8503 /rpt_family="Alu" 9070..9387 /rpt_family="Alu"
repeat_region	complement(9740..9845) /rpt_family="MER2" complement(10640..11015) /rpt_family="Alu"
repeat_region	11950..12250 /rpt_family="Alu" 12057..12085 /note="(A)29"
repeat_region	/rpt_type=tandem /rpt_unit=A 12365..12645 /rpt_family="Alu" 13727..13750 /note="(AC)12"
repeat_region	/rpt_type=tandem /rpt_unit=CC 13783..14024 /rpt_family="L1"
repeat_region	14175..14470 /rpt_family="Alu" complement(14906..15259) /standard_name="possible repeat"
repeat_region	15300..15613 /rpt_family="Alu" 16671..16680 /note="(T)20"
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repeat_region	complement(16993..17085) /rpt_family="MR42" complement(17678..18276) /rpt_family="Alu"
repeat_region	19305..19583 /rpt_family="Alu" 19914..19945

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  /rpt_family="Alu"
repeat_region
  21736, .22035
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  22017, .22038
  /note="(A)22038"
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  /rpt_family="Alu"
  23473, .23761
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repeat_region
  23744, .23767
  /note="(A)24"
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  /rpt_unit=A
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  complement(25727, .26471)
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  27774, .28057
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  28040, .28066
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  /rpt_unit=A
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  /rpt_family="MER20"
  complement(28769, .28838)
  /note="GRAIL 2 excellent exon, frame 0"
  28987, .29214
  /rpt_family="Alu"
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  29495, .29976
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  /note="100% identity EST OV84a10.x1"
  complement(30401, .30536)
  /note="GRAIL 2 excellent exon, frame 2"
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  complement(30682, .30733)
  /rpt_family="MIR"
  complement(31573, .31724)
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  complement(32159, .32232)
  /note="GRAIL 2 excellent exon, frame 2"
  32388, .32488
  /rpt_family="MLT1"
  32617, .32908
  /rpt_family="Alu"
  32977, .33088
  /rpt_family="MLT1"
  complement(33670, .33785)
  /rpt_family="Alu"
  complement(34021, .34144)
  /note="GRAIL 2 excellent exon, frame 2"
  complement(35238, .35331)
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40503, .40661,41868, .41972,42103, .42225,42492, .42569,
44379, .44507))
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/note="78%-100% protein identity GenPept:U18937"
complement(38069, .38215)
/note="GRAIL 2 excellent exon, frame 0"

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Best Local Similarity 16.7% Pred. No. 1,30e-03;
Matches 14; Conservative 45; Mismatches 24; Indels 1; Gaps 1;

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Db 15929 CMSRSKRGWGRYSKRYKCAAMATCKSS-KCWCSTYRMRKCYSCSYCCSSGKKYWC 15987
QY 2 CCGGAGCTCACTGCTCTCCATCCAGAGCGCAGTGCACATAGCGTCTGGCGTG 61
Db 15988 RCGMYTTCYYSKYKYSMSYCTC 16011
QY 62 CCCCTGCTCTCCTCTGACCTC 85

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RESULT 7
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DEFINITION Homo sapiens Chromosome 16 BAC clone CIT987SK-A-952P10, complete
sequence.
ACCESSION AC004787
NID 93337381
VERSION AC004787.1 GI:3337381
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Primates; Catarrhini; Hominoidea; Homo.

```

```

REFERENCE
AUTHORS Adams,M.D., Loftus,B.J., Zhou,L., Crosby,M., Fuhmann,J.,
Mason,T.M., Brandon,R., Kim,U.J., Kerlavage,A.R. and Venter,J.C.
Homo sapiens Chromosome 16 BAC clone CIT987SK-A-952P10
Unpublished
2 (bases 1 to 216021)
Adams,M.D. and Loftus,B.J.

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REFERENCE
AUTHORS Adams,M.D. and Loftus,B.J.
TITLE Direct Submission
JOURNAL Submitted (02-JUN-1998) The Institute for Genomic Research, 9712
Medical Center Dr., Rockville, MD 20850, USA, Email:
b.j.loftus@tigr.org
3 (bases 1 to 216021)
Adams,M.D. and Loftus,B.J.

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REFERENCE
AUTHORS Adams,M.D. and Loftus,B.J.
TITLE Direct Submission
JOURNAL Submitted (24-JUL-1998) The Institute for Genomic Research, 9712
Medical Center Dr., Rockville, MD 20850, USA
On Jul 24, 1998 this sequence version replaced gi:3241936.
Address all correspondence to: Mark Adams The Institute for Genomic
Research 9712 Medical Center Dr., Rockville, MD 20850, USA e-mail
sp6 end to T7 end. Genes were identified by a combination of five
methods including: XGRAIL (available by anonymous ftp from
athur.gem.cnl.gov), GeneFinder (Phil Green, University of
Washington), GenScan (Chris Burge,
http://gnomc.stanford.edu/~chris/GENSCAN.html) searches of the
complete sequence against a peptide database, and the Human gene
Index database at TIGR (http://www.tigr.org/db/hgi.html).
Genes without peptide homology having spliced EST hits are termed
'Unknown gene product'. Genes encoding tRNAs are predicted by
tRNAScan-SE (Sean Eddy, http://genome.wustl.edu/eddy/tRNAScan-SE/).

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BASE COUNT	60960 a 51778 c 49172 g 53987 t	124 others
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Db	1457 KXYMKSRRRARRSGSKKKKYYYYYYCYCYCYCYCYCGRAMMAAAAYYKRRCMAMY 1516	
Cp	222 TGAAGGAGATGGAGGAGGAGGCACACTTTCACGCAATTCCAGAGGTGACTCATAG 163	
Db	1517 YMRGRARKTYYYMARRGCARISRYKKKYAMNMYAMTWCAAAAAMAATYYNNMMYYKKW 1576	
Cp	162 GAAGAATTTTGCAAAAGAACCTCTTCAGCTTAGTTGCANAGTCATACC GGCCCTGT 103	
Db	1577 MYTCTCGAG 1585	
Cp	102 CCATGTGAG 94	
RESULT	8	
LOCUS	128278	215 bp DNA PAT 30-OCT-1996
DEFINITION	Sequence 5 from patent US 5569830.	
ACCESSION	128278	
NID	91819054	
VERSION	128278.1 GI:1819054	
KEYWORDS	.	
SOURCE	Unknown.	
ORGANISM	Unknown.	
REFERENCE	Unclassified.	
AUTHORS	1 (bases 1 to 215) Bennett,A., Labavitch,J.M., Powell,A. and Stetz,H.	
TITLE	Plant inhibitors of fungal polygalacturonases and their use to control fungal disease	
JOURNAL	Patent: US 5569830-A 5 29-OCT-1996;	
FEATURES	Location/Qualifiers	
source	1..215 /organism="unknown"	
BASE COUNT	15 a 8 c 25 g	26 t 141 others
ORIGIN		
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Db	18 CNDAKKDGGTTSWTCCCRHTWGCDITFTYVNNDSGHNKYSANANYNGGNNAK 77	
Cp	85 CCTGGCAGCTACATGCAACAGGCGCGGTGTGACTTTCACAACGTGAG-AAAGAGT 143	

Db	78	THYTTNTNAGDASTYTDSDYXNMGSSNGSDGDRSGADSDSSSTYTA	127
Ox	144	CTTTCTGTGCAAAATTCTCTCTATGAGATCCAGCTTCTCGGAATTCGTGAA	193
RESULT	9	AR024229	965 bp DNA PAT 04-DEC-1998
LOCUS	DEFINITION	Sequence 22 from patent US 5795961.	
ACCESSION	AR024229		
NID	93977523		
VERSION	AR024229.1	GI:3977523	
KEYWORDS	unknown.		
SOURCE	ORGANISM	unknown.	
REFERENCE	1 (bases 1 to 965)		
AUTHORS	Wallace,T.Paul, Harris,W.J., Carr,F.J., Old,L.J., Walt,S. and Kitamura,K.		
TITLE	Recombinant human anti-Lewis b antibodies		
JOURNAL	Patent: US 5795961-A 22 18-AUG-1998;		
FEATURES	Location/Qualifiers		
source	1..965		
BASE COUNT	192 a 170 c 226 g 205 t	172 others	
ORIGIN	/organism="unknown"		
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Best Local Similarity	15.6%; Freq. No. 1.90e-02;		
Matches	22; Conservative 59; Mismatches 60; Indels 0; Gaps 0;		
Db	790	GGVSTSTCTASDTTSTYMGVNRGRMGDYGGGYTNNGKRGRTVMTDTSNRSRSTVA	849
Cp	202	GGCAGACTTTTCAAGCAATTCACAGAGAGCTGACTCATAGGAGGAATTGTCCGAAAGA	143
Db	850	DPAVYVCVRGRSYSDSGDYGWGGTGTATVSHTYKDMTSSSSASVADRVTTCCRSSTHNG	909
Cp	142	CTCTCTTCAGCTGTAGTGTGAAGTCAATACCGGCCCTGTTCATGTGAGCTGCCAAGAG	83
Db	910	NTYWTYKGAKAIRYSNRGSVS 930	
Cp	82	GGTCAAGAGAGACAAAGGGG 62	
RESULT	10	MYB87256	1056 bp DNA MAM 02-JAN-1999
LOCUS	DEFINITION	Mustela vison GT dinucleotide repeat, chromosome 1q.	
ACCESSION	U87256		
NTD	94099442		
VERSION	U87256.1	GI:4099442	
KEYWORDS	American mink.		
SOURCE	ORGANISM	Mustela vison	
REFERENCE	1 (bases 1 to 1056)		
AUTHORS	Eukaryote; Metazoa; Chordata; Vertebrata; Mammalia; Euteria;		
TITLE	Carnivora; Fissipedia; Mustelidae; Mustela.		
JOURNAL	Submitted (27-JAN-1997) Breeding and Genetics, Danish Institute of		
FEATURES	Animal Science, Blichersøla K25, Tjele 8830, DK		
source	Location/Qualifiers		
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/db_xref="taxon:9667"			
/chromosome="1"			
/map="1q"			
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gagcattccacgtctgtgag"			
98..119			
/strand.name="1167F"			
complement(300..320)			
/strand.name="1167R"			
BASE COUNT	211 a 221 c 210 g 225 t	189 others	
ORIGIN			


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13147..13250,14614..14679,18199..18248,18465..18542,
18635..18746,23665..23755,24574..24709,37283..37493)
/notes="Hypothetical human protein (partial)"
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/evidence="not_experimental"
/product="R29828-1"
/protein_id="AAD03161.1"
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LMEKRLQVPEHLTLGQIVENWGNPQKESIRVPELIVTWHADNGSVKPC
KTDKALWLEKRLKMDSPILSSFEVTLLEHITNCVLVGHKATRLDGLCYSVK
MDNAEQFTTALRLYSLERINPDHSFVSCRAAFYRGLFEFGRIKAKKE
LRETLMSNAEDNRLTACSLVILGHIFVGLNHRNSNNVPAWQASKIPDSVL
WSSALLRDNKACGNADHQAQMHQNSQQLQDIEACSLPEHNLITVGVHWEG
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/notes="DSD similarity to overlapping ESTs: -AA963316
UI-R-EI-g1-c-06-0-UI-s1 UI-R-EI Rattus norvegicus CDNA
clone UI-R-EI-g1-c-06-0-UI 3'; (238..303); 88%
identity. -AA893275 EST197078 Normalized rat kidney; Bento
Soares Rattus sp. CDNA clone RKIB38 3' end; (258..323);
88% identity."
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clone UI-R-EI-g1-c-06-0-UI 3'; (304..399); 89%
identity. -AA893275 EST197078 Normalized rat kidney; Bento
Soares Rattus sp. CDNA clone RKIB38 3' end; (324..419);
89% identity. - (6480..6660) predicted exon, program:
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score: 76.000"
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frame: 0, quality: excellent, score: 94.000--DSD
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clone UI-R-EI-g1-c-06-0-UI 3'; (400..482); 88%
identity. - (8155..8240) AA893275 EST197078 Normalized rat
kidney; Bento Soares Rattus sp. CDNA clone RKIB38 3' end;
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frame: 1, quality: excellent, score: 100.000"
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frame: 0, quality: excellent, score: 100.000"
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frame: 2, quality: excellent, score: 97.000"
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19632..19928
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19929..20226
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Complement(20344..20655)
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Note: remainder of annotations omitted.
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Best Local Similarity 85.78; Pred. No. 2.76e-01;
Matches 30; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
Db 13976 TGTCCACGATGCGATCTGCGCTGCCCTTGCCCTC 14010
Gy 39 TGGCCACATGAGGCTGCGCTGCCCTTGCCCTC 73
RESULT 13
LOCUS A62989 60 bp DNA PAT 12-MAR-1998
DEFINITION Sequence 1 from Patent WO9720068.
ACCESSION A62989
NID 93716861
VERSION A62989.1 GI:3716861
KEYWORDS
SOURCE
ORGANISM
unclassified.
unclassified.
REFERENCE 1 (bases 1 to 60)
AUTHORS Cerum.H. and Seeger.C.
TITLE METHOD FOR GENERATING MULTIPLE DOUBLE STRANDED NUCLEIC ACIDS
JOURNAL Patent: WO 9720068-A 1 05-JUN-1997;
BOEHRINGER MANNHEIM GMBH (DE)
FEATURES
Location/Qualifiers
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/db_xref="taxon:32644"
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[illegible]

Query Match	13.58;	Score 31;	DB 12;	Length 114;
Best Local Similarity	4.68;	Pred. No. 5.71e-05;		
Matches	5;	Conservative 30;	Mismatches 74;	Indels 0;
			Gaps	0;

RESULT	8
ID	Q70468 standard; DNA; 114 BP.

FH	key	Location/Qualifiers
FT	misc_feature	55..60

FT /note- "this sequence represents 'Z'; Z can be a
FT sequence of 6, 9 or 12 nucleotides (see
FT comments)"

PN W054163187-A.
 PD 18-AUG-1994.
 PE 01-FEB-1993; U00977.
 PR 01-FEB-1993; US-013416.
 PR 30-DEC-1993; US-176500.
 PR 31-JAN-1994; US-189331.
 PA (DUNC-) UNIV. NORTH CAROLINA.
 PI Fowlkes DM, Kay BK.
 P1 MPI; 94-279739/34.
 DR P-PSDB; R65154.
 PT Identifying proteins or peptide(s) which bind a ligand - by
 PT screening a recombinant vector library expressing fusion proteins
 PT comprising a binding domain and an effector domain
 PS Disclosure: Page 33; 255pp: English.
 CC Q70466 is a generic DNA sequence used to generate random TSAR (Totally
 CC Synthetic Affinity Reagents) peptides. This generic formula can also be
 CC represented as follows: X(NNB)11(TCC)6X(NNB)7(TCC)(NNB)10Y. X
 CC and Y are flanking restriction sites (X is not the same as Y) that are
 CC not specified further. Other generic sequences are shown in Q70466-68.
 CC Other specific peptides generated by these generic sequences are shown in

Query Match	13.1%;	Score 30;	DB 12;	Length 114;
Best Local Similarity	3.8%;	Pred. No. 1.93e-04;		
Matches	4;	Conservative	30;	Mismatches 72;
			Indels	0;
			Gaps	0;

[illegible]

RESULT	9
ID	Q70470 standard; DNA; 114 BP

DT 10-Apr-1995 (first entry)
DE Generic DNA sequence to generate a random TSAR peptide library.
KW TSAR: totally synthetic affinity reagent; synthetic; binding domain
KW effector domain; concatenated heterocuticular protein; linker
KW direct; rapid; detection; screening; treatment; generic; ss.

FT	misc_feature	Location/Qualifiers
FT	FT	55..60
FT	FT	/*tag" a
FT	FT	/note "encoded by Z (see comments)"

PN MO9418318-R.
PD 18-AUG-1994.
PE 01-FEB-1994; U00977.
PR 01-FEB-1993; US-013416.
PR 30-DEC-1993; US-176500.
PR 31-JAN-1994; US-189331.
PA (UYNC-) UNIV NORTH CAROLINA.
PI Fowlkes DM, Kay BK;
MP; 94-279739/34.
DR P-PSDE: R58378.

PT Identifying proteins or peptide(s) which bind a ligand - by
PT screening a recombinant vector library expressing fusion proteins
PT comprising a binding domain and an effector domain
P5 Disclosure: Page 36: 253pp: English.
070410 is a generic DNA sequence used to generate random TSAR (totally
CC Synthetic Affinity Reagents) peptides This generic formula can also be
CC represented as follows: X(NNN)Y(CAC)(CAC)(NNB)8(CAC)6(CAC)8
CC as Y that are not specified further. The peptides generated by this and
CC as Y that are not specified further. The peptides generated by this and
CC other generic sequences (Q0471-73) have invariant histidine residues
CC incorporated into variant sequences. TSARs are concatenated
CC heterofunctional proteins or peptides, comprising at least two functional
CC regions - a binding domain with affinity for a ligand and a second
CC effector peptide portion that is chemically or biologically active. They
CC may further comprise a linker peptide between the 2 domains. The TSARs
CC or compounds comprising a TSAR binding domain can be used in vivo to
CC deliver a chemically or biologically active moiety, eg. metal ion,
CC radioisotope, peptide, toxin or enzyme, to the specific target or on the
CC cell. They can also replace the function of macromolecules, eg.

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(TM)

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MPsrch_mn n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 17:08:54 1999; Maspar time 20.47 Seconds

Tabular output not generated. 967.578 Million cell updates/sec

Title: >US-09-092-296-2

Description: (1-229) from US09092296.seq

Perfect Score: 229

N.A. Sequence: 1 ACCGGGACCTTCAGTGTCTCC.....CGATCTCCCTTCAGGACCA 229

Scoring table: TABLE default

Gap 6

Mmatch STD : Dbase 0; Query 0

Searched: 165359 seqs, 43243793 bases x 2

Post-processing: Minimum Match 08

Listing first 45 summaries

Database:

n-issued 1:5A_COMB 2:5B_COMB 3:5C_COMB 4:PCT9_COMB 5:backfile1

Statistics: Mean 7.362; Variance 4.165; scale 1.768

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description	Pred. No.
1	46	20.1	7218	2	US-08-232-Sequence 14, Applicat	3.37e-16
2	33	14.4	215	1	US-08-238-Sequence 5, Applicat	6.36e-08
3	31	13.5	965	3	US-08-388-Sequence 22, Applicat	1.05e-06
4	27	11.8	215	1	US-08-238-Sequence 5, Applicat	2.44e-04
5	27	11.8	965	3	PCT-US95-1-Sequence 22, Applicat	1.54e-01
6	22	9.6	75	4	PCT-US95-1-Sequence 98, Applicat	1.54e-01
7	22	9.6	82	4	PCT-US95-1-Sequence 97, Applicat	1.54e-01
8	22	9.6	3088	3	US-08-418-Sequence 145, Applicat	5.27e-01
9	22	9.6	65	1	US-08-471-Sequence 143, Applicat	5.27e-01
10	21	9.2	66	1	US-08-471-Sequence 143, Applicat	5.27e-01
11	21	9.2	68	1	US-08-471-Sequence 143, Applicat	5.27e-01
12	21	9.2	69	1	US-08-471-Sequence 143, Applicat	5.27e-01
13	21	9.2	74	4	PCT-US95-1-Sequence 100, Applicat	5.27e-01
14	21	9.2	74	4	PCT-US95-1-Sequence 99, Applicat	5.27e-01
15	21	9.2	75	4	PCT-US95-1-Sequence 98, Applicat	5.27e-01
16	21	9.2	81	4	PCT-US95-1-Sequence 97, Applicat	5.27e-01
17	21	9.2	81	4	PCT-US95-1-Sequence 96, Applicat	5.27e-01
18	21	9.2	82	4	PCT-US95-1-Sequence 95, Applicat	5.27e-01
19	21	9.2	92	3	US-08-333-Sequence 16, Applicat	5.27e-01
20	21	9.2	92	3	US-08-333-Sequence 16, Applicat	5.27e-01

21	21	9.2	906	3	US-09-031-Sequence 41, Applicat	5.27e-01
22	21	9.2	906	3	US-08-847-Sequence 41, Applicat	5.27e-01
23	21	9.2	906	3	US-08-847-Sequence 40, Applicat	5.27e-01
24	21	9.2	906	3	US-09-031-Sequence 40, Applicat	5.27e-01
25	21	9.2	906	3	US-09-031-Sequence 39, Applicat	5.27e-01
26	21	9.2	906	3	US-08-847-Sequence 39, Applicat	5.27e-01
27	21	9.2	906	3	US-08-847-Sequence 37, Applicat	5.27e-01
28	21	9.2	906	3	US-09-031-Sequence 37, Applicat	5.27e-01
29	21	9.2	906	3	US-09-031-Sequence 26, Applicat	5.27e-01
30	21	9.2	906	3	US-08-847-Sequence 26, Applicat	5.27e-01
31	21	9.2	909	3	US-09-031-Sequence 25, Applicat	5.27e-01
32	21	9.2	909	3	US-08-847-Sequence 25, Applicat	5.27e-01
33	21	9.2	911	3	US-09-031-Sequence 24, Applicat	5.27e-01
34	21	9.2	911	3	US-08-847-Sequence 22, Applicat	5.27e-01
35	21	9.2	911	3	US-08-847-Sequence 22, Applicat	5.27e-01
36	21	9.2	911	3	US-09-031-Sequence 36, Applicat	5.27e-01
37	21	9.2	5235	3	US-08-847-Sequence 36, Applicat	5.27e-01
38	21	9.2	5235	3	US-08-847-Sequence 35, Applicat	5.27e-01
39	21	9.2	5235	3	US-09-031-Sequence 35, Applicat	5.27e-01
40	21	9.2	5235	3	US-09-031-Sequence 34, Applicat	5.27e-01
41	21	9.2	5503	3	US-08-847-Sequence 34, Applicat	5.27e-01
42	21	9.2	5503	3	US-08-847-Sequence 34, Applicat	5.27e-01
43	21	9.2	5503	3	US-08-847-Sequence 32, Applicat	5.27e-01
44	21	9.2	5503	3	US-09-031-Sequence 32, Applicat	5.27e-01
45	20	8.7	66	4	PCT-US95-1-Sequence 93, Applicat	1.76e+00

ALIGNMENTS

RESULT 1
ID US-08-232-463-14 STANDARD, DNM, 7218 BP.
AC xxxxxx
DE Sequence 14, Application US/08232463
CC Patent No. 5670367
CC GENERAL INFORMATION:
CC APPLICANT: DORNER, F.
CC APPLICANT: SCHRIFFLER, F.
CC APPLICANT: FALKNER, F. G.
CC TITLE OF INVENTION: RECOMBINANT FOXP2 VIRUS
CC NUMBER OF SEQUENCES: 52
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Foley & Lardner
CC STREET: 1800 Diagonal Road, Suite 500
CC CITY: Alexandria
CC STATE: VA
CC COUNTRY: USA
CC ZIP: 22313-0299
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patentin release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/232,463
CC FILING DATE:
CC CLASSIFICATION: 435
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US/07/935,313.
CC FILING DATE:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: BENT, Stephen A.
CC REGISTRATION NUMBER: 29,768
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (703)836-9300
CC TELEFAX: (703)836-4109
CC TELE: 899149
CC INFORMATION FOR SEQ ID NO: 14:
CC SEQUENCE CHARACTERISTICS:

Query Match 13.5%; Score 31; DB 3; Length 965;

Best Local Similarity 19.3%; Pred. No. 1.05e-06;

Matches 23; Conservative 53; Mismatches 42; Indels 1; Gaps 1;

Db 845 SVTAADTAIVYCVNGSRYSDDGWDYGGTIVTVSSHVVDMTSSSSASVGDVRYTGRSST 904

19 CCTCCATCCGAGGAGCGAGTGCCATGCGGTGCGGCTGCGCCCTGCTCCTCTT 78

QY 905 TGNONTYWKAKYRVNSGYSRSGSGSDITTSDDTYTCGTHARTGCTVKG 963

79 GACC-CTCCTTGACGCTACATGGAACAGGCGCGGTATGACTTGCACTGAAGCTG 136

RESULT 4

ID US-08-238-163-5 STANDARD; DNA; UNC; 215 BP.

DT xxxxxx

DE Sequence 5, Application US/08238163

CC Sequence 5, Application US/08238163

CC Patent No. 5569830

CC GENERAL INFORMATION:

CC APPLICANT: BENNETT, Alan

CC APPLICANT: LABAVITCH, John W.

CC APPLICANT: POWELL, Ann

CC APPLICANT: STORZ, Henrik

CC TITLE OF INVENTION: PLANT INHIBITORS OF FUNGAL

CC TITLE OF INVENTION: POLYGALACTONASES AND THEIR USE TO CONTROL FUNGAL DISEASE

CC NUMBER OF SEQUENCES: 24

CC CORRESPONDENCE ADDRESSES:

CC ADDRESSEE: Townsend and Townsend Kourie and Crew

CC STREET: Stewart Street Tower, One Market Plaza

CC CITY: San Francisco

CC STATE: California

CC COUNTRY: US

CC ZIP: 94105-1493

CC COMPUTER READABLE FORM:

CC MEDIUM TYPE: Floppy disk

CC COMPUTER: IBM PC compatible

CC OPERATING SYSTEM: PC-DOS/MS-DOS

CC SOFTWARE: Patent Release #1.0, Version #1.25

CC CURRENT APPLICATION DATA:

CC APPLICATION NUMBER: US/08/238,163

CC FILING DATE: 03-MAY-1994

CC CLASSIFICATION: 800

CC ATTORNEY/AGENT INFORMATION:

CC NAME: Bastien, Kevin L.

CC REGISTRATION NUMBER: 34,774

CC REFERENCE/DOCKET NUMBER: 2307E-540

CC TELECOMMUNICATION INFORMATION:

CC TELEPHONE: (415) 543-9600

CC TELEFAX: (415) 543-5043

CC INFORMATION FOR SEQ ID NO: 5:

CC SEQUENCE CHARACTERISTICS:

CC LENGTH: 215 base pairs

CC TYPE: nucleic acid

CC STRANDEDNESS: Single

CC TOPOLOGY: unknown

CC MOLECULE TYPE: protein

CC FEATURE:

CC NAME/REV: misc.feature

CC LOCATION: 1..215

CC OTHER INFORMATION: /standard_name="Deduced amino acid

Query Match 11.8%; Score 27; DB 1; Length 215;
Best Local Similarity 20.0%; Pred. No. 2.44e-04;
Matches 22; Conservative 42; Mismatches 45; Indels 1; Gaps 1;

Db 18 CNDKAKDGNSTTSMTDCCNRTWGVCDTTRVYNDGNNKSSSNRYGNNVGA 77

OY 85 CCTTGACGCTACATGGAACAGGCGCGGTATGACTTGCACTGAAGCTG-AGGAGT 143

Db 78 THYTHTNVSGADSKTYTDSYASGTSSNGGTDGNNRSGADSYSSKRTAM 127

QY 144 CTTTCTGACAAATTCCTCTATGATGACGCTCCTGGAATTCCTTGA 193

RESULT 5

ID US-08-388-672A-22 STANDARD; DNA; UNC; 965 BP.

DT xxxxxx

DE Sequence 22, Application US/08388672A

CC Sequence 22, Application US/08388672A

CC Patent No. 5795961

CC GENERAL INFORMATION:

CC APPLICANT: Wallace, T. Paul

CC APPLICANT: Harris, William J.

CC APPLICANT: Carr, Frank J.

CC APPLICANT: Old, Lloyd J.

CC APPLICANT: Welt, Sydney

CC APPLICANT: Klamura, Kunio

CC TITLE OF INVENTION: Recombinant Human Anti-Lewis B

CC NUMBER OF SEQUENCES: 25

CC CORRESPONDENCE ADDRESSES:

CC ADDRESSEE: Felte and Lynch

CC STREET: 805 Third Avenue

CC CITY: New York

CC STATE: New York

CC COUNTRY: U.S.A.

CC ZIP: 10022

CC COMPUTER READABLE FORM:

CC MEDIUM TYPE: Floppy disk

CC COMPUTER: IBM PC compatible

CC OPERATING SYSTEM: PC-DOS/MS-DOS

CC SOFTWARE: Patent Release #1.0, Version #1.30

CC CURRENT APPLICATION DATA:

CC APPLICATION NUMBER: US/08/388,672A

CC FILING DATE: 14-FEB-1995

CC CLASSIFICATION:

CC ATTORNEY/AGENT INFORMATION:

CC NAME: Hanson, No. 5795961man D.

CC REGISTRATION NUMBER: 30,946

CC REFERENCE/DOCKET NUMBER: LUD 5409

CC TELECOMMUNICATION INFORMATION:

CC TELEPHONE: 212-688-9200

CC TELEFAX: 212-688-3884

CC INFORMATION FOR SEQ ID NO: 22:

CC SEQUENCE CHARACTERISTICS:

CC LENGTH: 965 base pairs

CC TYPE: nucleic acid

CC STRANDEDNESS: unknown

CC TOPOLOGY: unknown

CC MOLECULE TYPE: DNA (genomic)

CC SEQUENCE 965 BP; 192 A; 170 C; 226 G; 200 T; 177 OTHER.

Query Match 11.8%; Score 27; DB 3; Length 965;
Best Local Similarity 15.6%; Pred. No. 2.44e-04;
Matches 22; Conservative 59; Mismatches 60; Indels 0; Gaps 0;

Db 790 GGVNSTCTASDITTSYWGVRGKWDYGGGTNTNGRGVYTMADTSSNSSVTA 849

Cp 202 GCGAGACTTTTAAACAAATTCAGGAAGCTGACTCATAGAGAAATTCACAAAAG 143

Db 850 DTAIVYCVNGSRYSDDGWDYGGTIVTVSSHVVDMTSSASVGDVRYTGRSSTH 909

Cp 142 CTCCTGACGCTCATGCAAGATCATACCGCGCCCTGTCATGTGAGCTGCCAAGAG 83

Db 910 NTYVWKAKYRVNSGYSRSGSGSDITTSDDTYTCGTHARTGCTVKG 963

Cp 82 GGTAGAGAGACAAAGGG 62

RESULT 6

Mon Oct 25 11:53:51 1999

US-09-092-296-2.rni

Page 4

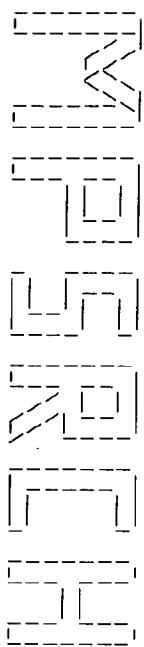
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ID      PCT-US95-11934-99 STANDARD; DNA; UNC; 75 BP.
DE      xxxxxx
DT
DB      Sequence 98, Application PC/TUS9511934
DY      Sequence 99, Application PC/TUS9511934
YY      GENERAL INFORMATION:
CC      APPLICANT: Cytogen Corporation
CC      TITLE OF INVENTION: Antigen Binding Peptides (Ablides) From
CC      TITLE OF INVENTION: Peptide Libraries
CC      NUMBER OF SEQUENCES: 103
CC      CORRESPONDENCE ADDRESS:
CC      ADDRESSEE: Pennie & Edmonds
CC      STREET: 1155 Avenue of the Americas
CC      CITY: New York
CC      STATE: New York
CC      COUNTRY: USA
CC      ZIP: 10036
CC      COMPUTER READABLE FORM:
CC      MEDIUM TYPE: Floppy disk
CC      COMPUTER: IBM PC Compatible
CC      OPERATING SYSTEM: PC-DOS/MS-DOS
CC      SOFTWARE: Patent Release #1.0, Version #1.30
CC      CURRENT APPLICATION DATA:
CC      APPLICATION NUMBER: PCT/US95/11934
CC      FILING DATE: 20-SEP-1995
CC      CLASSIFICATION:
CC      ATTORNEY/AGENT INFORMATION:
CC      NAME: Mistock, S. Leslie
CC      REGISTRATION NUMBER: 18,872
CC      REFERENCE/DOCKET NUMBER: 1101-196-228
CC      TELECOMMUNICATION INFORMATION:
CC      TELEPHONE: (212) 790-9090
CC      TELEFAX: (212) 869-9741/8864
CC      TELEX: 66141 PENNIE
CC      INFORMATION FOR SEQ ID NO: 99:
CC      SOURCE CHARACTERISTICS:
CC      LENGTH: 75 base pairs
CC      TYPE: nucleic acid
CC      STRANDEDNESS: single
CC      TOPOLOGY: linear
CC      MOLECULE TYPE: DNA (genomic)
CC      SEQUENCE 75 BP; 1 A; 1 C; 5 T; 61 OTHER.
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Query Match          9.6%; Score 22; DB 4; Length 75;
Best Local Similarity 8.7%; Pred.No.1,54e+01;
Matches    6; Conservative   19; Mismatches 44; Indels    0; Gaps    0.

Db         5 GNNBNNBNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNNBNN 64
           |
Dy         50 GGATGTCGGAGCGCCTTGTTCTTGCACCTCTTGAGCATGTGACACAGG 109
           :|||
Oy        110 CCGGGTAGTg 118

RESULT     7
ID      PCT-US95-11934-98 STANDARD; DNA; UNC; 81 BP.
DE      xxxxxx
DT
DB      Sequence 98, Application PC/TUS9511934
DY      Sequence 98, Application PC/TUS9511934
YY      GENERAL INFORMATION:
CC      APPLICANT: Cytogen Corporation
CC      TITLE OF INVENTION: Antigen Binding Peptides (Ablides) From
CC      TITLE OF INVENTION: Peptide Libraries
CC      NUMBER OF SEQUENCES: 103
CC      CORRESPONDENCE ADDRESS:
CC      ADDRESSEE: Pennie & Edmonds
CC      STREET: 1155 Avenue of the Americas
CC      CITY: New York
CC      STATE: New York
CC      COUNTRY: USA
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CC CC ZIP: 10036
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: PatentIn Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: PCT/US95/11934
CC FILING DATE: 20-SEP-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-196-228
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (212) 790-9090
CC TELEFAX: (212) 869-9741/8964
CC TELEX: 6614 PENNIE
CC INFORMATION FOR SEQ ID NO: 98:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 81 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: DNA (genomic)
CC SEQUENCE 81 BP: 6 A; 6 C; 4 G; 5 T; 60 OTHER.
S0
Query Match          9.68; Score 22; DB 4; Length 81;
Best Local Similarity 7.68; Fred. No. 1.54e-01;
Matches      5; Conservative      20; Mismatches 41; Indels    0; Gaps    0
Db       5 TGTCTAGANNVNNNVNNNVNNNVNNNVNNNVNNNVNNNVNNNVNNNVNNV 64
Cp       92 TGCCACGAGGGTCAAGAGGAGGACAGAGGGCGACCACCCATAGTGCGCATCTGC 33
Db       65 NNVVNV 70
Cp       32 TCCTGG 27
RESULT      8
ID PCT-US95-11934-97 STANDARD; DNA; UNC; 82 BP.
AC xxxxxx
DE Sequence 97, Application PC/TUS9511934
CC Sequence 97, Application PC/TUS9511934
CC GENERAL INFORMATION:
CC APPLICANT: Cyrogen Corporation
CC TITLE OF INVENTION: Antigen Binding Peptides (abtlides) From
CC NUMBER OF SEQUENCES: 103
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Pennie & Edmonds
CC STREET: 1155 Avenue of the Americas
CC CITY: New York
CC STATE: New York
CC COUNTRY: USA
CC ZIP: 10036
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: PatentIn Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: PCT/US95/11934
CC FILING DATE: 20-SEP-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Mistock, S. Leslie
CC REGISTRATION NUMBER: 18,872
CC REFERENCE/DOCKET NUMBER: 1101-196-228
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (212) 790-9090
```

(TM)

Release 3.1A John F. Collins, Biocomputing Research Unit.
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MSearch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 16:59:27 1999; MasPar time 446.22 Seconds

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Description: (1-229) from US09092296.seq

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N.A. Sequence: 1 ACCGGAGCTGACGTGCTCC.....CCATCTCCCTTCAGGAGCA 229

Comp: TGCCCTGAGGTACACAGAG.....GGTAGAGGAGGAGTCCCTGCT

Scoring table: TABLE default

Gap 6

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Searched: 2883791 seqs, 1171580779 bases x 2

Post-processing: Minimum Match 08

Listing first 45 summaries

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6:em-est9 7:em-gss1

Database:

genbank-est11

8:gb-est1 9:gb-est10 10:gb-est11 11:gb-est12 12:gb-est13

13:gb-est14 14:gb-est15 15:gb-est16 16:gb-est17

17:gb-est18 18:gb-est19 19:gb-est20 20:gb-est21

21:gb-est22 22:gb-est23 23:gb-est24 24:gb-est25

25:gb-est26 26:gb-est27 27:gb-est28 28:gb-est29

29:gb-est30 30:gb-est31 31:gb-est32 32:gb-est33 33:gb-est34

34:gb-est35 35:gb-est36 36:gb-est37 37:gb-est38 38:gb-est39

39:gb-est40 40:gb-est41 41:gb-est42 42:gb-est43

Statistics: Mean 9.755; Variance 2.089; scale 4.670

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,

and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description	Pred. No.
1	53	23.1	328	23	A1136523	UT-R-C2P-nq-e-02-0-UI	4.59e-50
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3	50	21.8	252	17	AA754459	97SN1787 Rice Immature	4.21e-45
4	40	17.5	247	17	AA754458	97SN1784 Rice Immature	4.11e-29
5	32	14.0	247	17	AA754458	97SN1784 Rice Immature	3.60e-17
6	32	14.0	2275	20	AF034173	AF034173 Human mRNA (T	3.60e-17
7	27	11.8	2275	20	AF034173	AF034173 Human mRNA (T	2.78e-10
8	24	10.5	348	19	F06958	HSC10C101. normalized 1	1.90e-06
9	23	10.0	238	12	AA376266	EST8815 HSC172 cells	3.12e-05
10	23	10.0	311	11	AA323964	EST26816 Cerebellum II.	3.12e-05

11	10.0	364	34	W33870	mc56b03.r1 Soares nous	3.12e-05
12	10.0	412	18	AA769782	ah71b05.s1 Soares_test	3.12e-05
13	10.0	426	8	T15255	cr5855_lambdAAPT Ric	3.12e-05
14	10.0	481	14	AA498488	vh44b06.r1 Barstead MP	3.12e-05
15	10.0	1287	20	AF038250	AF038250 Human mRNA (T	3.12e-05
16	9.6	308	8	DA0392	R1CS23424 Rice shoot O	4.75e-04
17	9.6	339	19	F08745	HSC1DB011 normalized 1	4.75e-04
18	9.6	395	41	A0375965	RPc111-161H11.TV RPc11	4.75e-04
19	9.6	404	17	AA730403	nv42c07.s1 NCI-CGAP_Ew	4.75e-04
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21	9.6	441	14	C28493	C28493 Rice callus CDN	4.75e-04
22	9.6	634	22	A1063013	G802423 Spine GH Dros	4.75e-04
23	9.6	1025	37	B12587	F20111-565.1 IGF Arabi	4.75e-04
24	9.2	299	8	M79528	we9700065 Mixed stage,	6.60e-03
25	9.2	370	25	A1128672	as91lne.r1 Neurospora	6.60e-03
26	9.2	384	21	A1160316	qb66d04.x1 Soares_feta	6.60e-03
27	9.2	401	42	A0444216	G85TC0313 Trypanosoma	6.60e-03
28	9.2	417	30	R36249	yh91f09.r1 Soares plac	6.60e-03
29	9.2	420	28	A1566143	tn53h10.x1 NCI-CGAP_K1	6.60e-03
30	9.2	422	10	AA275959	vc37e03.r1 Barstead MP	6.60e-03
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33	9.2	471	40	A0236981	RPc111-64172.TK RPc111	6.60e-03
34	9.2	494	20	AA898724	NCM6E877 Mycelial Neur	6.60e-03
35	9.2	499	37	B45020	HS-1060-B1-B01-MF.ab1	6.60e-03
36	9.2	502	15	AA586074	28723 Lambda-PRI2 Arab	6.60e-03
37	9.2	524	15	AA33967	mw24907.r1 Soares_mous	6.60e-03
38	9.2	533	15	AA593745	n183g11.s1 NCI-CGAP_Br	6.60e-03
39	9.2	550	20	AA898701	NCM6G777 Mycelial Neur	6.60e-03
40	9.2	567	30	R61539	yh16f01.s1 Soares_infa	6.60e-03
41	9.2	594	20	AA898768	NCM6C777 Perithecial N	6.60e-03
42	9.2	601	32	N37023	yy40d03.s1 Soares_mela	6.60e-03
43	9.2	688	37	AG015068	Homo sapiens genomic D	6.60e-03
44	9.2	699	36	AA140679	CK00462.5Spine CK Dros	6.60e-03
45	9.2	796	37	AF035986	Salmonella typhimurium	6.60e-03

ALIGNMENTS

RESULT	1	LOCUS	A1136523	328 bp	mRNA	EST	11-FEB-1999
DEFINITION	UT-R-C2P-nq-e-02-0-UI.s1	UT-R-C2P	Rattus norvegicus	CDNA clone			
ACCESSION	A1136523	UT-R-C2P-nq-e-02-0-UI 3'	mRNA sequence.				
VERSION	93637300						
KEYWORDS	A1136523.1	GI:3637300					
SOURCE	EST.						
ORGANISM	Norway rat.						
REFERENCE	Rattus norvegicus						
AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;						
TITLE	Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.						
JOURNAL	1 (bases 1 to 328)						
MEDLINE	Bonaldo,M.F., Lennon,G. and Soares,M.B.						
COMMENT	Normalization and subtraction: two approaches to facilitate gene						
	discovery						
	Genome Res. 5 (9), 791-806 (1996)						
	97044477						
	On Jan 14, 1998 this sequence version replaced gi:1877567.						

Contact: Soares, MB
Program for Rat Gene Discovery and Mapping
University of Iowa
451 Eckstein Medical Research Building Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: msoares@iuii.weeg.uiowa.edu
The sequence tag present in the cDNA between the NotI site and the
Oligo dT track served to identify it as a clone from the normalized
adult lung library. cDNA library Preparation: M. Fatima Bonaldo,
Ph.D. Clone distribution: clones will be available through Research
Genetics
Seq primer: M3 Forward.
Location/Qualifiers


```

/cultivar="M11veng23"
/notes="vector: pBluescript SK(+); Site_1: EcoRI; Site_2:
XhoI; Directional cDNA library inserted into lambda ZAP101
vector at 5' end with EcoRI and 3' end with Xho I site."
/db_xref="taxon:4530"
/map="6"
/clone="97SN1787"
/clone_lib="Rice Immature Seed Lambda ZAP101 cDNA Library"
/tissue_type="Immature Seed"
/dev_stage="5 days after pollination"
/lab_host="E. coli SOLR"

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	Best Local Similarity	9.8%	Pred. No.	4,21e+45	
	Matches	21	Conservative	108	Mismatches 83; Indels 2; Gaps 2
Db	22	SYBCHGNWVYCVASHGNTMSYNBCBTGCTDCDKANNSTWTGTGVMNBSGDHNYB	81		
Cp	224	CCGAGAGGACATGGACGAGCAGCAGCATTTTAAGCATTCCAGCAAGCGTAGACTAT	165		
Db	82	VBNITKYDNGNTPCSRPBYRYAAYHYDTMCBVNNYNNHHMBMYVBVTGTCOTMNC	141		
Cp	164	AGAGAGAAATTGTGCAAAAAGACTCCTTGACCTTCAGTTGCCAAATGCATACCCGGCCG	105		
Db	142	WBHYNTKCTAGSGHTSTINYKSSINTPTGYLBTIDSKMGY-C-SBYUKTHNKYSTPA	199		
Cp	104	TTCATGTAAGCTGCCCAAGAGGCTCACAAAGAGACAGAGGAGCACCAGACCCCATAG	45		
Db	200	TRSTCYCRKCYMMATKKVKTKHYVBGSBVD	233		
Cp	44	TGGCACTGCGCTCTCGGTGAGGAGACACT	11		

[illegible]

CONTACT: Eun M.Y.
 Department of Cytogenetics
 National Inst. of Agri. Sci. and Tech, RDA
 Suwon, Kyungido, Korea
 Tel: 62 331 290 0301
 Fax: 82 331 290 0307
 Email: myeun@sunsu.asti.re.kr
 Submitted by Baek Hie Nam, Dept of Biological Science, Myongji
 University, Yongin, Korea, 449-728 bhnahme@blosserver.myonji.ac.kr
Seq primer: M13 Reverse Primer.
Location/Qualifiers
 1. 247
SOURCE
 /organism="Oryza sativa"
 /cultivar="MilYang23"
 /note="Vector: pBluescript SK(+); Site_1: EcoRI; Site_2:
 XhoI; Directional cDNA library inserted into lambda ZAPIR
 vector at 5' end with EcoRI and 3' end with Xho I site."

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/map="6"
/clone="97SN1784"
/clone_lib="Rice Immature Seed Lambda ZAPII CDNA Library"
/tissue_type="Immature Seed"
/dev_stage="5 days after pollination"
/lab_host="E. coli SOLR"
BASE COUNT      7 a      16 c      21 g      34 t      169 others
ORIGIN
Query Match      17.5%; Score 40; DB 17; Length 247;
Best Local Similarity 10.7%; Pred. No. 4,11e-29;
Matches 22; Conservative 100; Mismatches 81; Indels 3; Gaps 3;

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[illegible]

LOCUS	5	247 bp	MRNA	EST	20-JAN-1998
DEFINITION	97SN1784 Rice Immature Seed Lambda ZAPRI cDNA Library Oryza sativa				
ACCESSION	CDNA clone 97SN1784, mRNA sequence.				
VERSION	92801154				
KEYWORDS	AA754458.1 GI:2801164				
SOURCE	EST.				
ORGANISM	Oryza sativa.				
REFERENCE	Oryza sativa				
AUTHORS	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Oryza.				
	1 (bases 1 to 247)				
	Nahn,B.H., Kim,J.K., Cheong,J.J., Kim,S.I., Hahn,T.R., Moon,E.P., Kim,W.Y., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y., Lee,M.C. and Eun,M.Y.				
	Large-scale Sequencing Analysis of ESTs from Rice Immature Seed				
	disrupted (1999)				
COMMENT	On Jan 14, 1998 this sequence version replaced gi:1797455.				

```

Contact: Eun M.Y.
Department of Cyrogenetics
National Inst. of Agri. Sci. and Tech, RDA
Suwon, Kyungido, Korea
Tel: 82 331 290 0301
Fax: 82 331 290 0307
Email: myeun@suncn0.asti.re.kr
Submitted by Baek Hie Nahm, Dept of Biological Science, Myongji
University, Yongin, Korea. 449-728 bhna@bioserver.myongji.ac.kr
Seq primer: M3 Reverse Primer.

FEATURES
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            /cultivar="Milyang23"
            /note="vector: pBluescript SK(+); site_1: EcoRI; site_2:
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            vector at 5' end with EcoRI and 3' end with Xho I site."
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            /map="6"
            /clone="g7snt1784"
            /clone_lib="Rice Immature Seed Lambda ZAPRII cDNA library"

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[illegible]

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LOCUS       7          AF034173      2275 bp      mRNA      EST      30-MAR-1998
DEFINITION  AF034173 Human mRNA (Tripodis and Ragousis) Homo sapiens cDNA
ACCESSION   AF034173
NID         92707735
VERSION     AF034173.1 GI:2707735
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (bases 1 to 348)
Auffray,C., Behar,G., Bols,F., Bouchier,C., da Silva,C.,
Davignes,M.D., Duprat,S., Houligatte,R., Jumeau,M.N., Lamy,B.,
Lorenzo,F., Mitchell,H., Mariage-Samson,R., Pletiot,G., Pouliot,Y.,
Sebastiant,Kabakthchis,C. and Tessier,A.
IMAGE: molecular integration of the analysis of the human genome
and its expression
C. R. Acad. Sci. III, Sci. Vie 318 (2), 263-272 (1995)
95277534
On Sep 21, 1992 this sequence version replaced gi:279286.

JOURNAL
COMMENT

REFERENCE
AUTHORS
TITLE

JOURNAL
COMMENT

On Jan 19, 1998 this sequence version replaced gi:2045115.

Contact: Tripodis, Nikos
Division of Medical and Molecular Genetics
Guys Hospital
7th floor, Guy's Tower, London SE1 9RT, UK
Email: nikos@hki.nl.
Location/Qualifiers
1..2275
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="6p21.3"
/clone="nticon2 contig"
/clone_1b="Human mRNA (Tripodis and Ragousis)"
BASE COUNT  438 a 619 c 470 g 599 t 149 others
ORIGIN
Query Match 11.8%; Score 27; DB 20; Length 2275;
Best Local Similarity 16.2%; Pred. No. 2,78e-10;
Matches 11; Conservative 39; Mismatches 17; Indels 1; Gaps 1
Db 1556 YWVCWCCTSKKASACAMRMKCMYMS-RSSSTVWGVMSGCGCTGATKRYKYSMTGWR 1614
QY 20 CTCGATCCGAGGCGCAGTGCGACTGTGGGCTCGGCGTCCCTCTCTCTCTG 79
Db 1615 WTWYMYNM 1622
QY : : : : :
80 ACCCTCCT 87

RESULT 8 F06958 348 bp mRNA EST 20-FEB-1995
LOCUS LOCUS
DEFINITION HSC10C101 normalized infant brain cDNA Homo sapiens cDNA clone
c-19c10, mRNA sequence.
ACCESSION F06958
NID 9672595
VERSION F06958.1 GI:672595
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (bases 1 to 348)
Auffray,C., Behar,G., Bols,F., Bouchier,C., da Silva,C.,
Davignes,M.D., Duprat,S., Houligatte,R., Jumeau,M.N., Lamy,B.,
Lorenzo,F., Mitchell,H., Mariage-Samson,R., Pletiot,G., Pouliot,Y.,
Sebastiant,Kabakthchis,C. and Tessier,A.
IMAGE: molecular integration of the analysis of the human genome
and its expression
C. R. Acad. Sci. III, Sci. Vie 318 (2), 263-272 (1995)
95277534
On Sep 21, 1992 this sequence version replaced gi:279286.

```


Contact: Maria M/Mouse EST Project
WashU-HHMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:224283
Seq primer: E9P1mer
High quality sequence stop: 360

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FEATURES
source      Location/Qualifiers
            1..412
            /organism="Homo sapiens"
            /note="Vector: pT7TD-Pac (Pharmacia) with a modified
            polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA"
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QY 8 CTTCACTCTCTCTCCATCCAGAGCCGACTGCGCCTATGG 50

RESULT 15
LOCUS AF038250 1287 bp mRNA EST 30-MAR-1998
DEFINITION AF038250 Human mRNA (Tripodis and Ragoussis) Homo sapiens cDNA
clone ntcon9, mRNA sequence.
ACCESSION AF038250
NID 92815880
VERSION AF038250.1 GI:2815880
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Tripodis, N. and Ragoussis, J.
TITLE Generation of a transcription map in the region immediately
centromeric to human MHC across the 6p21.2-6p21.3 chromosomal
boundary
JOURNAL Unpublished (1997)
COMMENT On Jan 19, 1998 this sequence version replaced gi:2045085.

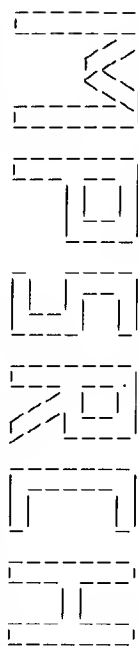
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/db_xref="taxon:9606"
/map="6p21.3"
/clone="ntcon9"
/clone_lib="Human mRNA (Tripodis and Ragoussis)"
Location/Qualifiers
1..1287
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="6p21.3"
/clone="ntcon9"
/clone_lib="Human mRNA (Tripodis and Ragoussis)"

BASE COUNT 349 a 219 c 293 g 361 t 65 others
ORIGIN

Query Match 10.0%; Score 23; DB 20; Length 1287;
Best Local Similarity 25.8%; Pred. No. 3.12e-05;
Matches 24; Conservative 32; Mismatches 37; Indels 0; Gaps 0;

DB 381 VVATCAGSCCVACVTCBCDCTGCGDTBHBVCKGCBANNDGATBCTCGKGVGA 440
QY 49 GGGCTGTGGGCTGCCCTTCTCTCTTGAACCTCTTGGCAGCTCAGTGAACAGG 108
DB 441 STCTHTNCCDCKTCGAGVTYNNHDMWMAAGA 473
QY 109 GCCGCGTATGACTTTCACACTGAAGCTGAAGA 141

Search completed: Sun Oct 24 17:07:02 1999
Job time : 455 secs.



(TM)

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MPearch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 16:22:19 1999: MasPar time 475.19 Seconds

Tabular output not generated. 1394.147 Million cell updates/sec

Title: >US-09-092-296-1
Description: (1-239) from US09092296.seq
Perfect Score: 238
N.A. Sequence: 1 GCCGACCGCGGACTTCAGTGT.....CCCTTCAGGACCAACGCTCA 239
Comp: CCGGTGGCCCTGAAAGTCACA.....GGCAAGTCCCTGGTGGCAGT

Scoring table: TABLE default
Gap 6

Match STD: Dbase 0; Query 0

Searched: 646147 seqs, 1385953633 bases x 2

Post-Processing: Minimum Match 0%

Listing first 45 summaries

Database:

emb158
1:em_ba1 2:em_ba2 3:em_fun 4:em_hlg 5:em_hum1 6:em_hum2
7:em_in 8:em_com 9:em_or 10:em_ov 11:em_pat 12:em_ph
13:em_pl 14:em_ro 15:em_sts 16:em_vi
genbank111

Database:
17:gb_ba1 18:gb_ba2 19:gb_hlg1 20:gb_hlg2 21:gb_in1
22:gb_in2 23:gb_com 24:gb_ov 25:gb_pat 26:gb_ph 27:gb_pl1
28:gb_pl2 29:gb_pl1 30:gb_pr2 31:gb_pr3 32:gb_ro
33:gb_st 34:gb_sts 35:gb_sy 36:gb_un 37:gb_vi

Statistics: Mean 9.607; Variance 4.824; scale 1.991

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query	Score	Match	Length	DB	ID	Description	Pred. No.
1	112	47.1	47323	31	AC005937	Homo sapiens clone UMG	7.72e-65	
2	47	19.7	7218	25	166494	Sequence 14 from paten	2.10e-15	
3	36	15.1	215	25	128278	Sequence 5 from patient	4.91e-08	
4	33	13.9	216021	31	HUAC004787	Homo sapiens Chromosom	4.91e-05	
5	31	13.0	965	25	AC024229	Sequence 22 from paten	8.69e-03	
6	29	12.2	74371	31	AC005369	Homo sapiens Chromosom	5.63e-03	
7	28	11.8	1056	23	WVU87256	Mustela vison GT dinuc	5.63e-03	
8	27	11.3	215	25	128278	Sequence 22 from patient	2.17e-02	
9	26	10.9	965	25	AR024229	Sequence 5 from paten	8.16e-01	
10	25	10.5	565	25	E04076	gDNA encoding envelope	3.00e-01	
11	25	10.5	60966	31	AC003030	Homo sapiens chromosome	3.00e-01	
12	25	10.1	60	25	A62989	Sequence 1 from Patient	1.07e+00	
13	24							

C	14	24	10.1	1056	23	WVU87256	Mustela vison GT dinuc	1.07e+00
C	15	24	10.1	175793	31	AC005920	Homo sapiens chromosome	1.07e+00
C	16	23	9.7	3390	29	HUMFUCAS	H.sapiens fucosidase P	3.72e+00
C	17	23	9.7	4472	17	NKSERBORF	Neisseria meningitidis	3.72e+00
C	18	23	9.7	133457	30	AC003939	Human PAC clone DU139	3.72e+00
C	19	23	9.7	192853	19	AC005096	Homo sapiens clone R33	3.72e+00
C	20	23	9.7	203418	19	AC004947	Homo sapiens clone DX1	3.72e+00
C	21	22	9.2	30	25	A62994	Sequence 6 from Patient	1.25e+01
C	22	22	9.2	108	21	D67227	Human (clone SY2/10) g	1.25e+01
C	23	22	9.2	1738	29	HUMGOLG1NB	D.melanogaster synapo	1.25e+01
C	24	22	9.2	2180	21	DROSTY	Human (clone SY2/10) g	1.25e+01
C	25	22	9.2	2273	32	MMBAEXN	Mus musculus Bak gene,	1.25e+01
C	26	22	9.2	3088	25	AR014574	Sequence 1 from patient	1.25e+01
C	27	22	9.2	3088	25	E10775	DNA encoding part of A	1.25e+01
C	28	22	9.2	3628	29	AB018292	Homo sapiens mRNA for	1.25e+01
C	29	22	9.2	3890	29	HSA010901	Homo sapiens KOC4 gene	1.25e+01
C	30	22	9.2	3514	32	AF090866	Mus musculus CDO (Cdo)	1.25e+01
C	31	22	9.2	6640	29	D63997	WORKING DRAFT SEQUENCE	1.25e+01
C	32	22	9.2	67919	20	AC006542	Homo sapiens mRNA for	1.25e+01
C	33	22	9.2	82098	31	AC006252	Homo sapiens 3p21.1 co	1.25e+01
C	34	22	9.2	124700	22	AC005558	Drosophila melanogaste	1.25e+01
C	35	22	9.2	135039	31	AC006060	Homo sapiens 3p22-8 PA	1.25e+01
C	36	22	9.2	151187	19	HS4608	Human DNA sequence ***	1.25e+01
C	37	22	9.2	175339	31	AC005772	Homo sapiens chromosome	1.25e+01
C	38	22	9.2	241407	20	AC003059	Mouse Chromosome 10 BA	1.25e+01
C	39	21	8.8	1755	29	HS297017	Homo sapiens mRNA for	4.06e+01
C	40	21	8.8	2817	27	AB012632	Robinia pseudocacacia 9	4.06e+01
C	41	21	8.8	3443	32	AB010145	Mus musculus gene for	4.06e+01
C	42	21	8.8	3247	32	AB010140	Mus musculus mRNA for	4.06e+01
C	43	21	8.8	46213	28	SPBC18H10	S.pombe chromosome II	4.06e+01
C	44	21	8.8	113887	20	AC006158	Homo sapiens clone NH0	4.06e+01
C	45	21	8.8	130560	19	HS508115	Human DNA sequence ***	4.06e+01

ALIGNMENTS

RESULT	LOCUS	1	AC005937	47323 bp	DNA	PRI	05-NOV-1998
DEFINITION	Homo sapiens clone UMG	C:370M23.002	from 6p21, complete sequence.				
ACCESSION	AC005937						
KEYWORDS	93845393						
VERSION	AC005937.1	GI:3845393					
SOURCE	HNG.						
ORGANISM	human.						
REFERENCE	Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;						
AUTHORS	Primates; Catarrhini; Hominiidae; Homo.						
TITLE	1 (bases 1 to 47323)						
REMARK	Janer,M., Guillaudoux,T., Vu,O., Kuttyavin,T., Harter,H. and						
	Geraghty,D.E.						
	Large scale sequence analysis of the human MHC class I region						
	Unpublished (1998)						
	Fred Hutchinson Cancer Research Center						
	The Clinical Research Division						
	1100 Fairview Ave. N., P.O. Box 19024						
	Seattle, WA 98109-1024						
	2 (bases 1 to 47323)						
	Submitted (05-NOV-1998) Human Genome Center, University of						
	Washington, Box 352145, Seattle, WA 98195, USA						
	University of Washington Human Genome Center						
	Box 352145 Seattle, WA 98195						
	Contact: Daniel E. Geraghty (geraghty@hccr.org)						
	Overlapping Sequences:						
	5': UMG:370M23.013 (Genbank Accession: AC005330)						
	3': UMG:Y67C112 (Genbank Accession: AC004211)						

Sequence Quality Assessment:
This entry has been annotated with sequence quality
estimates computed by the Phrap assembly program.
All manually edited bases have been reduced to quality zero.
Quality levels above 40 are expected to have less than


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note="GRAIL 2 excellent exon, frame 1"
36392..36663
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36801..37222
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/db_xref="dbEST:A1025011"
36901..37164
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39071..39205,39332..39630,39935..40048,40300..40410
40503..40661,41868..41972,42103..42225,42492..42569
44379..44507))
/standard_name="histidyl-tRNA synthetase"
/note="78%-100% protein identity GenPept:U18937"
complement(38069..38215)
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Note: remainder of annotations omitted.
Query Match
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Matches 14; Conservative 43; Mismatches 24; Indels 1; Gaps 1
Db 15929 CMSRSKSRKGNGVSMKTRICAMATMKSS-KCMWSYMRMKVCYSYCGSGKKYWC 15987
OY 6 CCAGGACTTCAGTGTCTCCCATCCAGGACGACGATGGCACTATGGTGTTGGGCTG 65
OY 66 CCCCTGTCTCCTCTGTAACCTC 89
FEATURES
source
MY087256 1056 bp DNA MAN 02-JAN-1999
Musstela vison GT dinucleotide repeat, chromosome 1q.
ND 94099442
U87256.1 GI:4099442
KEYWORDS
SOURCE
American vink.
Musstela vison.
ORGANISM
Eukaryote; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Canivora; Fissipedia; Mustelidae; Mustela.
1 (bases 1 to 1056)
Briegleb,R., Shkrti,N.M., Malchenko,S., Koroleva,I., and Lohi,O.
Direct Submission
Submitted (27-JAN-1997) Breeding and Genetics, Danish Institute o
Animal Science, Blichersalle K25, Tjele 8830, DK
location/Qualifiers
1..1056
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gaggaatctaccgttgttag"
98..119
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complement(300..320)
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BASE COUNT 211 a 221 c 210 g 225 t 189 others
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Best Local Similarity 12.8%; Pred. No.5,63e-03;
Matches 12; Conservative 49; Mismatches 33; Indels 0; Gaps 0;
Db 576 SCRHVBNMNMCKGSKCSKIGDKMSCAYCGRRRCRSRYMMNRQVSGSWARCC 635
OY 3 CCACGGAGCTAGTGTCTCCATCCAGGACGACGATGGCACTATGGTGTTGGGCTG 62

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QY	63	CTGCCCTTGTCTCCTCTGACCCCTCTGGCA	96
Db	636	CDDKSGDGNHCKSRKRKYKMDRHHKSKMA	669
LOCUS		: : : : : : : : : : : : : : :	
DEFINITION	8	HUAC004787 216021 bp DNA PRI	24-JUL-1998
ACCESSION		Homo sapiens Chromosome 16 BAC clone C119878K-A-952F10, complete sequence.	
NID		AC004787	
KEYWORDS		G3337381	
SOURCE		AC004787.1 GI:3337381	
ORGANISM		human.	
REFERENCE		Homo sapiens	
AUTHORS		Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;	
JOURNAL		Primates; Catarrhini; Hominoidea; Homo.	
TITLE		1 (bases 1 to 216021)	
REFERENCE		Adams,M.D., Loftus,B.J., Zhou,L., Crosby,M., Fuhrmann,J.,	
AUTHORS		Mason,T.M., Brandon,R., Kim,U.T., Kerlavage,A.R. and Ventner,J.C.	
JOURNAL		Homo sapiens Chromosome 16 BAC clone C119878K-A-952F10	
TITLE		Unpublished	
REFERENCE		2 (bases 1 to 216021)	
AUTHORS		Adams,M.D. and Loftus,B.J.	
JOURNAL		Direct Submission	
TITLE		Submitted (02-JUN-1998) The Institute for Genomic Research, 9712	
REFERENCE		Medical Center Dr, Rockville, MD 20850, USA, Email:	
AUTHORS		bjloftus@tigr.org	
JOURNAL		3 (bases 1 to 216021)	
TITLE		Adams,M.D. and Loftus,B.J.	
REFERENCE		Direct Submission	
AUTHORS		Submitted (24-JUL-1998) The Institute for Genomic Research, 9712	
JOURNAL		Medical Center Dr., Rockville, MD 20850, USA	
TITLE		On JUL 24, 1998 this sequence version replaced gi:3241936.	
REFERENCE		Address all correspondence to: Mark Adams The Institute for Genomics	
AUTHORS		Research 9112 Medical Center Dr, Rockville, MD 20850, USA e-mail	
JOURNAL		address: bjloftus@tigr.org. The orientation of the sequence is from the	
TITLE		sfp end to 3' end. Genes were identified by a combination of five	
REFERENCE		methods including: xcrail (available by anonymous ftp from	
AUTHORS		archur.epm.ornl.gov), GeneFinder (Phil Green, University of	
JOURNAL		Washington). Genscan (Chris Burge,	
TITLE		http://genomic.stanford.edu/~chris/GENSCANW.html) searches of the	
REFERENCE		complete sequence against a peptide database, and the Human gene	
AUTHORS		Index database at TIGR (http://www.tigr.org/cdb/hg1.htm).	
JOURNAL		Genes without peptide homology having spliced EST hits are termed	
TITLE		'Unknown gene product'. Genes encoding tRNAs are predicted by	
REFERENCE		tRNAscan-SE (Sean Eddy, http://genome.wustl.edu/eddy/tRNAscan-SE/).	
AUTHORS		Location:Qualifiers	
JOURNAL		1..216021	
TITLE		/organism="Homo sapiens"	
REFERENCE		/db_xref="taxon:9606"	
AUTHORS		/chromosome="16"	
JOURNAL		/map="116q21.22"	
TITLE		/clone="A-952F10"	
REFERENCE		21765..21872	
AUTHORS		/note="7766, STS1-CSRL-2793-uA/CSRL-2793-uZ, Chr. -, Homo	
JOURNAL		sapiens"	
TITLE		/db_xref=dbSTS:G02280"	
REFERENCE		73826..73943	
AUTHORS		/note="7608, STS1-CSRL-2491-uA/CSRL-2491-uZ, Chr. -, Homo	
JOURNAL		sapiens"	
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AUTHORS		/note="16084, CHLC.GCT10B02, Chr. -, Homo sapiens"	
JOURNAL		/db_xref="dbSTS:G09703"	
TITLE		175810..175945	
REFERENCE		/note="16316, CHLC.GCT15C04, Chr. -, Homo sapiens"	
AUTHORS		/db_xref="dbSTS:G09935"	
JOURNAL		199463..199572	
TITLE		/note="9824, MT-3555, Chr. 16, Homo sapiens"	
REFERENCE		/db_xref="dbSTS:G04338"	
AUTHORS		60960 a 51778 c 49172 g 53987 t 124 others	


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LKQIQCIQDTSTLHDELLPSNPADLFHMIKREHMCVLYLVATVHNSQGLERAO
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clone UI-R-E1-g1-c-06-0-UI-3'; (238..303); 88%
identity. --AA893275 EST197078 Normalized rat kidney, Bento
Scores Rattus sp. CDNA clone R1B38 3' end; (258..323);
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complement(5602..5683)
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clone UI-R-E1-g1-c-06-0-UI-3'; (304..399); 89%
identity. --AA893275 EST197078 Normalized rat kidney, Bento
Scores Rattus sp. CDNA clone R1B38 3' end; (324..419);
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frame: 0, quality: excellent, score: 94.000--BDS
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clone UI-R-E1-g1-c-06-0-UI-3'; (400..482); 88%
identity. --(8155..8240) AA893275 EST197078 Normalized rat
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19925..20226
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Note: remainder of annotations omitted.
Query Match 10 5% Score 25; DB 31; Length 60966;
Best Local Similarity 85.7%; Pred. No. 3.00e-01;
Matches 30; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
DB 13976 TGTCCAGCATGATGGCGTGGCCCTTGCCTC 14010
Gy 43 TGGCCATCATGCGGTCTGGCGTCCCTGTCTCTC 77
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RESULT	13
LOCUS	A62989 60 bp DNA
DEFINITION	Sequence 1 from Patent WO9720068.
ACCESSION	A62989
NID	93716861
VERSION	A62989.1 GI:3716861
KEYWORDS	unidentified.
SOURCE	unidentified.
ORGANISM	unclassified.
REFERENCE	1 (bases 1 to 60)
AUTHORS	Oertum,H. and Seeger,C.
TITLE	METHOD FOR GENERATING MULTIPLE DOUBLE STRANDED NUCLEIC ACIDS
JOURNAL	Patient: WO 9720068-A 1 05-JUN-1997;
BOEHRINGER MANNHEIM GMBH (DE)	
FEATURES	Location/Qualifiers
source	1..60
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repeat_region /rpt_family="GC-rich" complement(19845, .19992)
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repeat_region /rpt_family="(CAA)n" complement(20648, .20927)
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repeat_region /rpt_family="MIR" complement(21690, .21999)
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repeat_region /rpt_family="AluJo" complement(25393, .25453)
repeat_region /rpt_family="L2" complement(25736, .25870)
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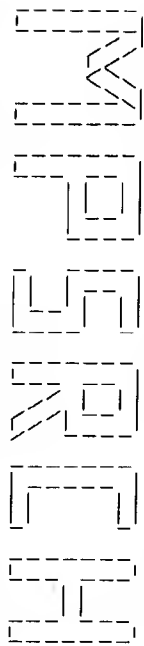
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Note: remainder of annotations omitted.

Query Match 10.1%; Score 24; DB 31; Length 175793;
Best Local Similarity 72.3%; Pred. No. 1.07e+00;
Matches 47; Conservative 0; Mismatches 17; Indels 1; Gaps 1;

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DB 18839 GGCAGCCAGACCCAGGCGTCACTGCGCCAGGAGGCGTG-CAACAGAGCTTG 18897
CP 67 GGCAGCCAGACCCAGGCGTCACTGCGCCAGGAGGCGTGCAATGCC 8
DB 18898 GGTGC 18902
CP 7 GGTGC 3
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Search completed: Sun Oct 24 16:30:24 1999
Job time : 485 secs.



(TM)

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Msearch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 16:38:53 1999; Maspar time 73.89 Seconds

Tabular output not generated. 693.101 Million cell updates/sec

Title: >US-09-092-296-1
Description: (1-239) from US09092296.seq
Perfect score: 238
N.A. Sequence: 1 GCCCAGCGGACTGTACGT.....CCCTTCAGGACGACGCTCA 239
Comp: CCGGTGGCCCTGATGACAC.....GGGAAGTCCCTGTGCGCAGT

Scoring table: TABLE default
Gap 6

Mmatch STD : Dbase 0; Query 0

Searched: 271905 seqs, 107135622 bases x 2

Post-processing: Minimum Match 08
Listing first 45 summaries

Database: n-geneseq35

1:part1 2:part2 3:part3 4:part4 5:part5 6:part6 7:part7
8:part8 9:part9 10:part10 11:part11 12:part12 13:part13
14:part14 15:part15 16:part16 17:part17 18:part18
19:part19 20:part20 21:part21 22:part22 23:part23
24:part24 25:part25 26:part26 27:part27 28:part28
29:part29 30:part30 31:part31 32:part32 33:part33
34:part34 35:part35 36:part36 37:part37 38:part38
39:part39 40:part40 41:part41 42:part42 43:part43
44:part44 45:part45 46:part46 47:part47 48:part48
49:part49 50:part50 51:part51 52:part52 53:part53
54:part54 55:part55 56:part56 57:part57 58:part58
59:part59 60:part60

Statistics: Mean 7.797; Variance 4.949; scale 1.575

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description	Pred. No.
1	232	97.5	439	60	V84366 Human stomach carcinoma	1.08e-132
2	37	13.5	91	9	Q51746 Oligonucleotide probe	3.39e-08
3	37	13.5	204	1	N81164 Base substituted E.co	3.39e-08
4	37	15.5	204	1	N81164 Base substituted E.co	3.39e-08
5	36	15.1	91	9	Q51746 Oligonucleotide probe	1.21e-07
6	31	13.0	114	12	Q70470 Generic DNA sequence	6.14e-05
7	31	13.0	114	12	Q70467 Generic DNA sequence	6.14e-05
8	30	12.6	114	12	Q70468 Generic DNA sequence	2.07e-04
9	30	12.6	114	12	Q70470 Generic DNA sequence	2.07e-04

C	10	30	12.6	114	12	Q70468	Generic DNA sequence	2.07e-04
C	11	29	12.2	114	12	Q70467	Generic DNA sequence	6.89e-04
C	12	29	12.2	114	12	Q70465	Generic DNA sequence	6.89e-04
C	13	29	12.2	114	12	Q70469	Generic DNA sequence	6.89e-04
C	14	28	11.8	114	12	Q70465	Generic DNA sequence	2.27e-03
C	15	28	11.8	178	32	T76405	Human endothelin-1 an	2.27e-03
C	16	27	11.3	114	12	Q70466	Generic DNA sequence	7.35e-03
C	17	27	11.3	172	32	T76363	Human interleukin 8 a	2.35e-02
C	18	26	10.9	114	12	Q70472	Generic DNA sequence	2.35e-02
C	19	26	10.9	114	12	Q70469	Generic DNA sequence	7.40e-02
C	20	25	10.5	39	7	Q51787	Mixed oligonucleotide	7.40e-02
C	21	25	10.5	114	12	Q70472	Substance P antisense	7.40e-02
C	22	25	10.5	250	32	T76438	Generic DNA sequence	2.29e-01
C	23	24	10.1	114	12	Q70473	Generic DNA sequence	2.29e-01
C	24	24	10.1	114	12	Q70473	Generic DNA sequence	2.29e-01
C	25	24	10.1	114	12	Q70473	Generic DNA sequence	6.97e-01
C	26	23	9.7	114	12	Q70471	Generic DNA sequence	6.97e-01
C	27	23	9.7	114	12	Q70471	Generic DNA sequence	6.97e-01
C	28	23	9.7	565	6	Q35072	HCV envelope region n	6.97e-01
C	29	22	9.2	36	2	Q11195	Ballast Constituent c	2.08e+00
C	30	22	9.2	75	21	T13612	DC43 TSAR library gen	2.08e+00
C	31	22	9.2	82	21	T13611	DC43 TSAR library gen	2.08e+00
C	32	22	9.2	82	21	T13610	DC43 TSAR library gen	2.08e+00
C	33	22	9.2	89	32	T76219	Human IL5 antisense o	2.08e+00
C	34	22	9.2	91	46	V44650	Mammalian DNA replica	2.08e+00
C	35	22	9.2	190	32	T76452	Chymase antisense oli	2.08e+00
C	36	22	9.2	264	32	T76445	Substance P receptor	2.08e+00
C	37	22	9.2	3088	16	T05628	ADP ribosylation fact	2.08e+00
C	38	21	8.8	168	32	T76270	Human MNCF antisense	6.06e+00
C	39	21	8.8	908	51	V63025	B. malayi ankyrin cDN	6.06e+00
C	40	21	8.8	908	52	V63315	Nucleotide nbnan908	6.06e+00
C	41	21	8.8	908	51	V63024	B. malayi ankyrin nm	6.06e+00
C	42	21	8.8	909	51	V63014	D. immitis ankyrin nd	6.06e+00
C	43	21	8.8	911	52	V63312	Nucleotide ndlan911	6.06e+00
C	44	21	8.8	911	51	V63012	D. immitis ankyrin nd	6.06e+00
C	45	21	8.8	5503	51	V63020	D. immitis ankyrin nd	6.06e+00

ALIGNMENTS

RESULT 1
ID V84366; standard; cDNA to mRNA; 439 BP.
AC V84366;
DI 30-MAR-1999 (first entry)
DE Human stomach carcinoma cDNA clone HP10408.
KW Transmembrane protein; HP10408; human; stomach cancer; ds.
OS Homo sapiens.
FH Key
FT Location/Qualifiers
FT 75..311
FT /tag= a
FT /note= "cDNA comprising the coding region (minus
the stop codon) is claimed (Claim 3)"
DR W0985508-A2.
DR 10-DEC-1998.
DR J02445.
DR 03-JUN-1998; J02445.
DR 03-JUN-1997; JP-14948.
PR (PROT-) PROTEGENE INC.
PA (SAG) SAGAMI CHEM RES CENTRE.
PI Kato S, Sekine S, Yamaguchi T;
DR WPI, 99-045730/04.
DR P-PDB; W88498.
FT New human proteins containing transmembrane domains and their
FT encoding sequences - useful in the preparation of antibodies and
FT large-scale protein production, gene diagnosis, and gene therapy
PS Claim 4; Page 15; 178bp; English.
CC This is the nucleotide sequence of cDNA clone HP10408, which
CC includes a coding region (also claimed) for a novel human
CC transmembrane protein (see W88498). The clone was isolated from a
CC stomach cancer cDNA library using a signal sequence detection
CC method, and by protein synthesis by in vitro translation. The
CC encoded protein has a putative signal sequence and a putative
CC internal transmembrane domain. The invention provides nucleotide
CC sequences (see W84359-76) coding for 18 transmembrane proteins


```
CC variable 3' ends generated in this way are used as primers for
CC reverse transcriptase. Nucleotides are misincorporated by the
CC transcriptase and the molecules are completed to forms that can be
CC amplified and then expressed in a suitable host-vector system.
CC The sequence covers all 176 diff base substitutions, most of which
CC occurred singularly in any given mutant.
CC See also P80575.
SQ Sequence 204 BP; 21 A; 47 C; 17 G; 11 T; 108 Others;

Query Match
Best Local Similarity 15.5% Score 37; DB 1; Length 204;
Matches 8; Conservative 54; Mismatches 39; Indels 0; Gaps 0;

Db 86 ymcttthyrrdnvrygynrsdaawyccyrsvydcymachdhbyvbbyyny 145
::| : : : : : : : : | : | : : : : : : : : : : : : : : : : :
Cp 172 TCATGAGCAAGCATTTGTGCACAAAGAAGCTTCACGTTCAGTTCGAAGTCATACC GGCG 113
::| : : : : : : : : | : | : : : : : : : : : : : : : : : : :
Db 146 hhmnmncgcchmhvhcnhvbmhmhnwaryvhdrrdyhc 186
::| : : : : : : : : | : | : : : : : : : : : : : : : : : : :
Cp 112 CCTGTTCCATGTGGAGCTGCCAAGAGGGGTCCAAGAGGAGAC 72

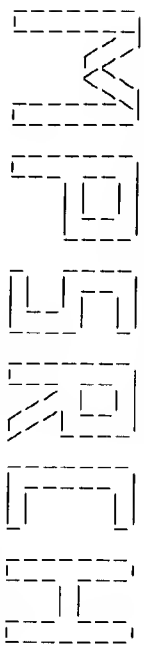
RESULT
ID 051746 standard; cDNA; 91 BP.
AC 051746;
DT 31-MAY-1994 (first entry)
DE Oligonucleotide probe MK14-A
KW Oligonucleotide; DNA probe; mycobacteria; disease diagnosis;
OS Synthetic.
PN EP-571911-A.
PD 01-DEC-1993.
PF 24-MAY-1993; 108325.
PR 25-MAY-1992; US-889651.
PA (BECT ) BECTON DICKINSON CO.
PI Shank DD, Spears PA;
DR WPI: 93-378844/48.
PT New oligo:nucleotide probes specific for Mycobacteria - used for
PT detection and amplification of Mycobacteria nucleic acid in
PT samples
PS Claim 3, Page 14; 23pp; English.
CC Oligonucleotide probe MK14-A consists of nucleotides 5-95 of MK14
CC (O51135). It hybridized to all spp. of mycobacteria tested, but
CC cross reacted to a few non-mycobacterial spp. The probe may
CC be useful as an initial screen for mycobacterial infection.
CC See also O51735-45 and O51147-59.
CC Sequence 91 BP; 5 A; 17 C; 15 G; 4 T;
SQ

Query Match
Best Local Similarity 15.1% Score 36; DB 9; Length 91;
Matches 0; Conservative 43; Mismatches 7; Indels 0; Gaps 0;

Db 11 ssahyvvhvshhsbvhybhvybvhhvvbhvybhvyrvs 60
::| : : : : : : : : | : | : : : : : : : : : : : : : : : : :
XY 7 CGGACTCAGCTGCTCCTCCATCCAGAGCCAGCTGGCCACTAGTGgg 56

RESULT
ID 070470 standard; DNA; 114 BP.
AC 070470;
DT 10-APR-1995 (first entry)
DE Generic DNA sequence to generate a random TSAR peptide library.
KW TSAR: totally synthetic affinity reagent; synthetic; binding domain;
KW effector domain; concatenated heterofunctional protein; linker;
KW direct; rapid; detection; screening; treatment; generic; ss.
OS Synthetic.
FT key Location/Qualifiers
FH misc-feature 53..60
FT /tag= a
FN MO94I8318-A.
PN 18-AUG-1994.
FE 01-FEB-1994; U00977.
```

[illegible]



(TM)

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MPerch_nm n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 16:40:29 1999; MasPar time 21.17 Seconds

Tabular output not generated. 976,635 Million cell updates/sec

Title: >US-09-092-296-1

Description: (1-239) from US09092296.seq

Perfect Score: 238

N.A. Sequence:

Comp: 1 GGCCAGCGGAGCTTAGTGT.....CCCTTACAGGACACGAGCTCA 239
CCGGTGGCCCTGAAATGACAC.....GGGAATGCTCCGTGTCGACG

Scoring table: TABLE default

Gap 6

Nmatch STD : Dbase 0; Query 0

Searched: 165359 seqs, 43243793 bases x 2

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database:

n-issued

1:5A.COMB 2:5B.COMB 3:5C.COMB 4:PCT9.COMB 5:backfiles1

Statistics: Mean 7.391; Variance 4.188; scale 1.765

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description	Pred. No.
1	47	19.7	7218	2 US-08-332-	Sequence 14, Applicati	8.84e-17
2	36	15.1	215	1 US-08-238-	Sequence 5, Applicatio	1.01e-09
3	31	13.0	965	3 US-08-388-	Sequence 22, Applicati	1.18e-06
4	27	11.3	215	1 US-08-238-	Sequence 2, Applicatio	2.69e-04
5	26	10.9	965	3 US-08-388-	Sequence 22, Applicati	1.01e-03
6	22	9.2	75	4 PCT-US95-1	Sequence 99, Applicati	1.67e-01
7	22	9.2	81	4 PCT-US95-1	Sequence 97, Applicati	1.67e-01
8	22	9.2	82	4 PCT-US95-1	Sequence 97, Applicati	1.67e-01
9	22	9.2	3088	3 US-08-418-	Sequence 1, Applicatio	1.67e-01
10	21	8.8	65	1 US-08-471-	Sequence 144, Applicat	5.69e-01
11	21	8.8	66	1 US-08-471-	Sequence 144, Applicat	5.69e-01
12	21	8.8	68	1 US-08-471-	Sequence 143, Applicat	5.69e-01
13	21	8.8	69	1 US-08-471-	Sequence 143, Applicat	5.69e-01
14	21	8.8	74	4 PCT-US95-1	Sequence 100, Applicat	5.69e-01
15	21	8.8	74	4 PCT-US95-1	Sequence 98, Applicati	5.69e-01
16	21	8.8	75	4 PCT-US95-1	Sequence 92, Applicati	5.69e-01
17	21	8.8	81	4 PCT-US95-1	Sequence 97, Applicati	5.69e-01
18	21	8.8	82	4 PCT-US95-1	Sequence 97, Applicati	5.69e-01
19	21	8.8	82	4 PCT-US95-1	Sequence 97, Applicati	5.69e-01
20	21	8.8	906	3 US-09-031-	Sequence 41, Applicati	5.69e-01

21	21	8.8	906	3 US-08-847-	Sequence 41, Applicati	5.69e-01
22	21	8.8	906	3 US-08-847-	Sequence 40, Applicati	5.69e-01
23	21	8.8	906	3 US-09-031-	Sequence 39, Applicati	5.69e-01
24	21	8.8	908	3 US-09-031-	Sequence 39, Applicati	5.69e-01
25	21	8.8	908	3 US-08-847-	Sequence 37, Applicati	5.69e-01
26	21	8.8	908	3 US-08-847-	Sequence 37, Applicati	5.69e-01
27	21	8.8	908	3 US-09-031-	Sequence 26, Applicati	5.69e-01
28	21	8.8	909	3 US-08-847-	Sequence 25, Applicati	5.69e-01
29	21	8.8	909	3 US-09-031-	Sequence 25, Applicati	5.69e-01
30	21	8.8	909	3 US-09-031-	Sequence 25, Applicati	5.69e-01
31	21	8.8	909	3 US-08-847-	Sequence 24, Applicati	5.69e-01
32	21	8.8	911	3 US-08-847-	Sequence 24, Applicati	5.69e-01
33	21	8.8	911	3 US-08-847-	Sequence 22, Applicati	5.69e-01
34	21	8.8	911	3 US-09-031-	Sequence 36, Applicati	5.69e-01
35	21	8.8	911	3 US-09-031-	Sequence 36, Applicati	5.69e-01
36	21	8.8	5235	3 US-08-847-	Sequence 35, Applicati	5.69e-01
37	21	8.8	5235	3 US-09-031-	Sequence 35, Applicati	5.69e-01
38	21	8.8	5235	3 US-08-847-	Sequence 34, Applicati	5.69e-01
39	21	8.8	5503	3 US-08-847-	Sequence 32, Applicati	5.69e-01
40	21	8.8	5503	3 US-09-031-	Sequence 32, Applicati	5.69e-01
41	21	8.8	5503	3 US-09-031-	Sequence 93, Applicati	1.89e+00
42	21	8.8	5503	3 US-08-847-	Sequence 93, Applicati	1.89e+00
43	21	8.8	5503	3 US-09-031-	Sequence 93, Applicati	1.89e+00
44	20	8.4	66	4 PCT-US95-1	Sequence 5, Applicatio	1.89e+00
45	20	8.4	1185	1 US-08-049-	Sequence 5, Applicatio	1.89e+00

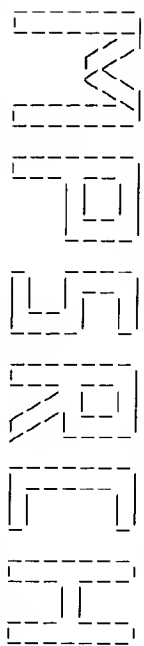
ALIGNMENTS

RESULT 1
ID US-08-232-463-14 STANDARD: DNA: UNC: 7218 BP.
AC xxxxxx

DE Sequence 14, Application US/08232463
CC Patent No. 5670367
CC GENERAL INFORMATION:
CC APPLICANT: DORNER, F.
CC APPLICANT: SCHNEIFLINGER, F.
CC APPLICANT: FALKNER, F. G.
CC TITLE OF INVENTION: RECOMBINANT FOXP2 VIRUS
CC NUMBER OF SEQUENCES: 32
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Foley & Lardner
CC STREET: 1800 Diagonal Road, Suite 500
CC CITY: Alexandria
CC STATE: VA
CC COUNTRY: USA
CC ZIP: 22313-0299
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: PatentIn Release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/232.463
CC FILING DATE:
CC CLASSIFICATION: 435
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US/07/935.313
CC FILING DATE:
CC APPLICATION NUMBER: EP 91 114 300.6
CC FILING DATE: 26-AUG-1991
CC ATTORNEY/AGENT INFORMATION:
CC NAME: BENT, Stephen A.
CC REGISTRATION NUMBER: 29,768
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (703)836-9300
CC TELEFAX: (703)683-4109
CC TELEX: 899149
CC INFORMATION FOR SEQ ID NO: 14:
CC SEQUENCE CHARACTERISTICS:

CC STRANDEDNESS: unknown
CC TOPOLOGY: unknown
CC MOLECULE TYPE: DNA (genomic)
SQ SEQUENCE 965 BP; 192 A; 170 C; 226 G; 200 T; 177 OTHER.
Query Match 13.0%; Score 31; DB 3; Length 965;
Best Local Similarity 19.3%; Pred. No. 1.18e-06;
Matches 23; Conservative 53; Mismatches 42; Indels 1; Gaps 1;
Db 845 SVTADIVYVYVGRSVDSDGDTGTTVYSVVVDMTSSSSASVDRVTTTGST 904
QY 23 CTTCCATCCAGAGGCGCAGTGCCTGAGGCTGCGGCTGCTGCTGCTCTT 82
Db 905 TEGNGTYWTKGAKRYVSNRSVSGSGSDTYTSSDATYTCGTHARTGTGK 963
QY 83 GACC-CTCCTTGACGCTCAGCATGAGACGGCCGGGTATGACTTGCACCTGACACTG 140
RESULT 4
ID US-08-238-163-5 STANDARD; DNA; UNC; 215 BP.
AC xxxxxx
DE Sequence 5, Application US/08238163
CC Sequence 5, Application US/08238163
CC Patent No. 5569830
CC GENERAL INFORMATION:
CC APPLICANT: BENNETT, Alan
CC APPLICANT: LABAVITCH, John M.
CC APPLICANT: POWELL, Ann
CC APPLICANT: STOTZ, Henrik
CC TITLE OF INVENTION: PLANT INHIBITORS OF FUNGAL
CC TITLE OF INVENTION: POLYGALACTURONASES AND THEIR USE TO CONTROL FUNGAL DISEASE
CC NUMBER OF SEQUENCES: 24
CC CORRESPONDENCE ADDRESSES:
CC ADDRESSEE: Townsend and Townsend Kourie and Crew
CC STREET: Stewart Street Tower, One Market Plaza
CC CITY: San Francisco
CC STATE: California
CC COUNTRY: US
CC ZIP: 94105-1493
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patent In Release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/238,163
CC FILING DATE: 03-MAY-1994
CC CLASSIFICATION: 800
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Bastian, Kevin L.
CC REGISTRATION NUMBER: 34,774
CC REFERENCE/DOCKET NUMBER: 2307E-540
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (415) 543-9600
CC TELEFAX: (415) 543-5043
CC INFORMATION FOR SEQ ID NO: 5:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 215 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: single
CC TOPOLOGY: unknown
CC MOLECULE TYPE: protein
CC FEATURE:
CC NAME/KEY: misc_feature
CC LOCATION: 1..215
CC OTHER INFORMATION: /standard_name="Deduced amino acid
CC OTHER INFORMATION: sequence of Pgip from bean."
SQ SEQUENCE 215 BP; 15 A; 8 C; 25 G; 26 T; 141 OTHER.
Query Match 11.3%; Score 27; DB 1; Length 215;
Best Local Similarity 20.0%; Pred. No. 2.69e-04;
Matches 22; Conservative 42; Mismatches 45; Indels 1; Gaps 1;

Db 18 CNDKAKIDGNTTSSWTTDCCNRGTGCDITDITRVNNDGHNKYSANYGNVGA 77
QY 89 CTTGCAGCTCAGCATGAGACGGCGGGGTATGACTTTCGACTGACACTG-AGAGCT 147
Db 78 THYTHHNVGSDSKTYVDSYNAGTSSNGCTDGNRSGADSYGSKTAM 127
QY 148 CTTTCTGACAAATTCCTCTATGATGATGACGCTCTTGAAATTCCTTGA 197
RESULT 5
ID US-08-388-672A-22 STANDARD; DNA; UNC; 965 BP.
AC xxxxxx
DE Sequence 22, Application US/08388672A
CC Sequence 22, Application US/08388672A
CC Patent No. 5795961
CC GENERAL INFORMATION:
CC APPLICANT: Wallace, T. Paul
CC APPLICANT: Harris, William J.
CC APPLICANT: Carr, Frank J.
CC APPLICANT: Old, Lloyd J.
CC APPLICANT: Welt, Sydney
CC APPLICANT: Kitamura, Kunio
CC TITLE OF INVENTION: Recombinant Human Anti-Lewis B
CC NUMBER OF SEQUENCES: 25
CC CORRESPONDENCE ADDRESSES:
CC ADDRESSEE: Felfe and Lynch
CC STREET: 805 Third Avenue
CC CITY: New York
CC STATE: New York
CC COUNTRY: U.S.A.
CC ZIP: 10022
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patent In Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/388,672A
CC FILING DATE: 14-FEB-1995
CC CLASSIFICATION:
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Hanson, No. 5795961man D.
CC REGISTRATION NUMBER: 30,946
CC REFERENCE/DOCKET NUMBER: LUD 5409
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: 212-688-9200
CC TELEFAX: 212-838-3884
CC INFORMATION FOR SEQ ID NO: 22:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 965 base pairs
CC TYPE: nucleic acid
CC STRANDEDNESS: unknown
CC TOPOLOGY: unknown
CC MOLECULE TYPE: DNA (genomic)
SQ SEQUENCE 965 BP; 192 A; 170 C; 226 G; 200 T; 177 OTHER.
Query Match 10.9%; Score 26; DB 3; Length 965;
Best Local Similarity 17.8%; Pred. No. 1.01e-03;
Matches 16; Conservative 43; Mismatches 30; Indels 1; Gaps 1;
Db 781 GURHUVHVGAVRSTCTASDTTYSYWGWRGRGWGDTGGVTVNKGRRVTMDTSS 840
CP 124 GTCATACCGCGGCTGT-TCCATGTGAGCTGCCAAGAGGTTCAGAGAGAGAGG 66
Db 841 NSRSVTADIVYVYVGRSVDSDGDTW 870
CP 65 CAGCCAGACCCCATAGTGGCAGCTGCGCT 36
RESULT 6



(TM)

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MPearch_n n.a. - n.a. database search, using Smith-Waterman algorithm

Run on: Sun Oct 24 16:30:43 1999; MasPar time 462.56 Seconds

Tabular output not generated. 1210.689 Million cell updates/sec

Title: >US-09-092-296-1
Description: (1-239) from US09092296.seq
Perfect Score: 238
N.A. Sequence: 1 GCCGCGGCGGAGCTTCAGTGT.....CCCTTCAGGACGACGCTCA 239
Comp: CCGGTCGCGGCTTGAAGTACACA.....GGGAACTCCCTGTGCGCAGT

Scoring table: TABLE default
Gap 6

Mmatch STD: Dbase 0; Query 0

Searched: 2883791 segs, 1171580779 bases x 2

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database:

embl-est58
1:em-est10 2:em-est11 3:em-est17 4:em-est18 5:em-est12
6:em-est9 7:em-gss1
genbank-est111

8:gb-est1 9:gb-est10 10:gb-est11 11:gb-est12 12:gb-est13
13:gb-est14 14:gb-est15 15:gb-est16 16:gb-est17
17:gb-est18 18:gb-est19 19:gb-est20 20:gb-est21
21:gb-est22 22:gb-est23 23:gb-est24 24:gb-est25
25:gb-est26 26:gb-est27 27:gb-est28 28:gb-est29
29:gb-est30 30:gb-est31 31:gb-est32 32:gb-est33 33:gb-est34
34:gb-est35 35:gb-est36 36:gb-est37 37:gb-gss1 38:gb-gss2
39:gb-gss3 40:gb-gss4 41:gb-gss5 42:gb-gss6

Statistics: Mean 9.792; Variance 2.100; scale 4.663

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query	Score	Match	Length	DB	ID	Description	Pred. No.
1	57	23.9	252	17	AA75459	97SN1787	Rice Immature	1.50e-56
2	54	22.7	328	23	AT136523	UT-R-C2P-ng-e-02-0-UT.	1.62e-51	
3	51	21.4	252	17	AA75459	97SN1787	Rice Immature	1.52e-46
4	33	16.4	247	17	AA754458	97SN1784	Rice Immature	2.04e-27
5	36	13.0	2275	20	AF034173	AF034173	Human mRNA (T	1.76e-18
6	31	13.0	2275	20	AF034173	AF034173	Human mRNA (T	1.24e-15
7	27	11.3	2275	20	AF034173	AF034173	Human mRNA (T	3.49e-10
8	24	10.1	348	19	F06958	HSC10C101	normalized i	2.31e-06
9	23	9.7	238	12	AA376266	EST88915	HSC172 cells	3.76e-05
10	23	9.7	311	11	AA323964	EST26816	Cerebellum IT	3.76e-05

C	11	23	9.7	364	34	W33870	mc56b03.r1 Soares	3.76e-05
C	12	23	9.7	426	18	AA769782	ah11b05.s1 Soares	3.76e-05
C	13	23	9.7	412	8	T15255	cr5855 lambdaZAPST	3.76e-05
C	14	23	9.7	481	14	AA498488	vh44b06.r1 Barstead	3.76e-05
C	15	23	9.7	1287	20	AF038250	AF038250 Human mRNA	3.76e-05
C	16	22	9.2	308	8	D40392	R1C2242A Rice shoot	3.65e-04
C	17	22	9.2	339	19	F08745	HSC1D501 normalized	5.65e-04
C	18	22	9.2	404	17	AA730403	hw42c07.s1 NC1.CGAP	5.65e-04
C	19	22	9.2	433	34	W79098	z075h10.r1 Soares	5.65e-04
C	20	22	9.2	441	14	C28493	C28493 Rice callus	5.65e-04
C	21	22	9.2	634	22	A1063013	GH02423.5prime GH	5.65e-04
C	22	22	9.2	1025	37	B12587	E22011-Sp6.1 1GF Arabi	5.65e-04
C	23	21	8.8	288	11	AA335414	EST39832 Epdidiymus	7.76e-03
C	24	21	8.8	299	8	M79528	WES700065 Mixed stage	7.76e-03
C	25	21	8.8	317	19	F10052	HSC39H122 normalized	7.76e-03
C	26	21	8.8	343	8	T47417	YD11F12.r1 Stratagene	7.76e-03
C	27	21	8.8	384	23	A1160316	qb66d04.x1 Soares	7.76e-03
C	28	21	8.8	395	41	A0375965	RPC111-161H11.TV RPI1	7.76e-03
C	29	21	8.8	401	42	A0444216	GSTC0313 Trypanosoma	7.76e-03
C	30	21	8.8	424	33	W13114	ma66e10.r1 Soares	7.76e-03
C	31	21	8.8	429	28	A1509530	vx09h04.r1 Soares	7.76e-03
C	32	21	8.8	440	18	AA822673	vw44f01.r1 Soares	7.76e-03
C	33	21	8.8	468	35	AA046996	zfs0a10.r1 Soares	7.76e-03
C	34	21	8.8	479	26	A1388157	MCSP4817 Subtracted	7.76e-03
C	35	21	8.8	499	37	B43020	HS-1060-B1-B01-MF	7.76e-03
C	36	21	8.8	516	31	H41536	Yp11c01.s1 Soares	7.76e-03
C	37	21	8.8	524	10	AA239967	mw24g07.r1 Soares	7.76e-03
C	38	21	8.8	550	20	AA898701	NCM6G77 Mycelial	7.76e-03
C	39	21	8.8	601	32	R61539	Yh16f01.s1 Soares	7.76e-03
C	40	21	8.8	637	23	N37023	Yy40d03.s1 Soares	7.76e-03
C	41	21	8.8	688	37	AG015068	Homo sapiens genomic	7.76e-03
C	42	21	8.8	694	36	AA140679	CSRL-133c3-u CSRL	7.76e-03
C	43	21	8.8	749	37	B01542	12711-Sp.1 TAMU Arabi	7.76e-03
C	44	21	8.8	762	37	B19344	Salmonella typhimurium	7.76e-03
C	45	21	8.8	796	37	AF035986		

ALIGNMENTS

RESULT	1	AA75459	252 bp	mRNA	EST	20-JAN-1998
LOCUS		97SN1787	Rice Immature seed	lambda ZAPIT	CDNA LIBRARY	Oryza sativa
DEFINITION		CDNA clone 97SN1787, mRNA sequence.				
ACCESSION		AA754459				
VERSION		92801165				
KEYWORDS		AA754459.1	GI:2801165			
SOURCE		EST.				
ORGANISM		Oryza sativa.				
		Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Oryza.				
REFERENCE		1 (bases 1 to 252)				
AUTHORS		Nahm,B.H., Kim,J.K., Cheong,J.U., Kim,S.I., Hahn,T.R., Moon,E.P., Kim,W.T., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y., Lee,M.C. and Eun,M.Y.				
TITLE		Large-scale Sequencing Analysis of ESTs from Rice Immature Seed				
JOURNAL		Unpublished (1998)				
COMMENT		On Jan 14, 1998 this sequence version replaced gi:1797457.				

Contact: Eun M.Y.
Department of Cytogenetics
National Inst. of Agri. Sci. and Tech, RDA
Suwon, Kyunggido, Korea
Tel: 82 331 290 0301
Fax: 82 331 290 0307
Email: myeun@sunsu20.asi.re.kr
Submitted by Baek Hie Nahm, Dept of Biological Science, Myongji
University, Yongin, Korea, 449-728 binahmed@server.myongji.ac.kr
Seq primer: M13 Reverse Primer.
Location/Qualifiers
1..252
/organism="Oryza sativa"

Query Match	23.9%	Score 57	DB 17	Length 252
Best Local Similarity	11.4%	Pred. No. 1.50e-56		
Matches	24	Conservative	109	Mismatches 75; Indels 3; Gaps 3

RESULT	2		
LOCUS	A1136523	328 bp	mRNA
DEFINITION	UI-R-C2P-ng-e-02-0-UI.s1	UI-R-C2P	Rattus norvegicus cDNA clone
	UI-R-C2P-ng-e-02-0-UI 3'		mRNA sequence.

ACCEPTATION
 A1136923
 NID
 93637800
 VERSION
 A1136923.1 GI:3637300
 KEYWORDS
 EST.
 SOURCE
 Norway rat.
 ORGANISM
 Rattus norvegicus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 REFERENCE
 1 (bases 1 to 328)
 AUTHORS
 Donaldson,M.F., Lennon,G. and Soares,M.B.
 TITLE
 Normalization and subtraction: two approaches to facilitate gene
 discovery
 JOURNAL
 Genome Res. 6 (3), 791-806 (1996)
 MEDLINE
 97044477
 COMMENT
 On Jan 14, 1998 this sequence version replaced gi:1877567.

FEATURES
source
 Contact: Soares, MB
 Program for Rat Gene Discovery and Mapping
 University of Iowa
 451 Eckstein Medical Research Building Iowa City, IA 52242, USA
 Tel: 319 335 8250
 Fax: 319 335 9565
 Email: mscoates@blue.veeg.uiowa.edu
 The sequence tag present in the cDNA between the NotI site and the
 oligo-OT track served to identify it as a clone from the normalized
 adult lung library, cDNA Library Preparation: M. Fatima Bonaldi,
 Ph.D. Clone distribution: clones will be available through Research
 Genetics
 Seq primer: M13 Forward.
 Location/Qualifiers:
 1..328
 /organism="Rattus norvegicus"
 /strain="Sprague-Dawley"
 /note="Vector: pT7T3D-Pac (Pharmacia) with a modified

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Query Match      72.2%, Score 54, Db 23, Length 328,
Best Local Similarity 24.7%, Prid. No. 1, 62e-51,
Matches 89, Conservative 0, Mismatches 30, Indels 1, Gaps 1

Db 181 TGACCTGTCGCCCGCTGAGAGGTGAGAGGAGAGCATCTTTTGAGCATGTCAGAA 240
Cp 239 TAGCCGTGCTCCCTCAAGAGCATGAGAGGAGAGGAGAGN-TTTTCAGCATTCAGGA 181
Db 241 AGCCGGAGCTTGGAGAGCTTGGCCCTGATAGCTTCGTCACTTTACTTCGAAGTCA 300
Cp 180 AGCTGAGCATATGAGAGGAATTTGTGAGAAAAGCTCTTACGCTTAGTTCAGAAAGTCA 121

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LOCUS	AA754459	252 bp	mRNA	EST	20-JAN-1998
DEFINITION	cDNA clone 97SN1/87 Rice Immature seed Lambda ZAPRI cDNA Library Oryza sativa				
ACCESSION	AA754459				
NID	92801165				
VERSION	AA754459.1	GI:2801165			
KEYWORDS	EST.				
SOURCE	Oryza sativa.				
ORGANISM	Oryza sativa				
REFERENCE	Eukaryota: Vitellidplantae; Streptophyta; Embryophyta; Tracheophyta; eumhyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Oryza.				
AUTHORS	1. (bases 1 to 252) Nahn,B.H., Kim,J.K., Cheong,J.J., Kim,S.I., Nahn,T.R., Moon,E.P., Kim,T.H., Kim,W.Y., Yang,M.S., Park,R.D., Sohn,U.I., Kang,K.Y., Lee,M.C. and Eun,M.Y.				
TITLE	Large-Scale Sequencing Analysis of ESTs from Rice Immature Seed				
JOURNAL	Unpublished (1998)				
COMMENT	On Jan 14, 1998 this sequence version replaced gi:1797457.				

Contact: Eun M.Y.
Department of Cytogenetics
National Inst. of Agri. Sci. and Tech, RDA
Suwon, Kyunggi-do, Korea
Tel: 82 331 290 0301
Fax: 82 331 290 0307
Email: myeun@sunsu20.asi.re.kr
Submitted by Baek Hie Nahm, Dept of Biological Science, Myongji
University, Yongin, Korea, 449-728 binaim@bserver.myongji.ac.kr

	BASE COUNT	7 a	16 c	21 g	34 t	169 others
	ORIGIN					
	Query Match	13.9%	Score 33;	D8 17;	Length 247;	
	Best Local Similarity	13.3%	Pred. No. 1,76e-18;			
	Matches	26;	Conservative	88;	Mismatches 79;	Indels 3; Gaps 3;
		/db_xref="taxon:4530"				
		/map="6"				
		/clone="97SN1784"				
		/clone_lib="Rice Immature Seed Lambda ZAPII cDNA Library"				
		/tissue_type="Immature Seed"				
		/dev_stage="5 days after pollination"				
		/lab_host="E. coli SOLR"				
Db	48	GTTTTCNDSDNAHCRITYBMYARS-KYCYGTBYTSNNDDTGTGCTGYKTYVNVHSW	106			
Oy	10	GACTCATAGTCCTCCATCCACAGAGCGCAGTGCCACATAGGGGTGTGGCGGCCCC	69			
Db	107	NNRCNSNYSVVMBETAYCDVBHYDBRAHWVDTRCTNDRGYCNTASDNGTSATKRVTGYD	166			
Oy	70	TTCGTCTCTCTCTTGACCCTCCCTTGCGACGCACACATGAGAAGCGGGCGGT-ATGACCTTG	128			
Db	167	KTDSDCGGCGCKRYVYGS-SBYBRGVNVWRTSMTDKSTAKBSMDRSRRYHGRV	225			
Oy	129	CACATGAAGCTGAAAGAGCTTTCTCACAAATTCCTCTATGATGCAGCTTCOTGGAA	188			
Db	226	MBNRKRGSKRNMTDTK	241			
Oy	189	TTCGTTGMAAANTCTG	204			
RESULT	6	AF034173	2275 bp	mRNA	EST	30-MAR-1998
LOCUS		AF034173	Human mRNA (Tripodis and Ragousais)	Homo sapiens CMA		
DEFINITION			Clone ntcon2 contig.	mRNA sequence.		
ACCESSION		AF034173				
NID		g2707735				
VERSION		AF034173.1	GI:2707735			
KEYWORDS		EST.				
SOURCE		human.				
ORGANISM		Homo sapiens				
		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;				
		Eutheria; Primates; Catarrhini; Hominoidea; Homo.				
REFERENCE		1 (bases 1 to 2275)				
AUTHORS		Tripodis,N. and Ragousais,J.				
TITLE		Generation of a transcription map in the region immediately centromeric to human HMC across the 6p21.2-6p21.3 chromosomal boundary				
JOURNAL		Unpublished (1997)				
COMMENT		On Jan 19, 1998 this sequence version replaced gi:2045115.				
		Contact: Tripodis, Nikos				
		Division of Medical and Molecular Genetics				
		Guys Hospital				
		7th floor, Guy's Tower, London SE1 9RT, UK				
		Email: nikose@kcl.nl				
FEATURES		Location/Qualifiers				
source		1..2275				
		/organism="Homo sapiens"				
		/db_xref="taxon:9606"				
		/map="6p21.3"				
		/clone_ntcon2 contig"				
		/clone_lib="Human mRNA (Tripodis and Ragousais)"				
BASE COUNT	438 a	619 c	470 g	599 t	149 others	
ORIGIN						
Query Match	13.0%	Score 31;	DB 20;	Length 2275;		
Best Local Similarity	9.6%;	Pred. No. 1.24e-15;				
Matches	11;	Conservative	64;	Mismatches 39;	Indels 1;	Gaps 1;
Db	1480	RKKRKRRKKRKKRRKTKRRKYRAMMAACAAMACAMTWYKMKMGKKCKTKRKKKTTS	1539			
Oy	217	ATTGAGGAGGAGGAGGAGGCAATTTCACCAATTCACGAGACCTGCACTCATPAGAGAAATTT	158			

RESULT	7	AF034173	2275 bp	mRNA	EST	30-MAR-1998
LOCUS	AF034173	Human mRNA (Tripodis and Ragoussis)	Human sapiens	CDNA		
DEFINITION	clone ntcon2 contig, mRNA sequence.					
ACCESSION	AF034173					
NID	92707735					
VERSION	AF034173.1	GI:2707735				
KEYWORDS	EST.					
SOURCE	human.					
ORGANISM	human sapiens					
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;					
	Eutheria; Primates; Catarrhini; Hominoidea; Homo.					
REFERENCE	1 (bases 1 to 2275)					
AUTHORS	Tripodis, N. and Ragoussis, J.					
TITLE	Generation of a transcription map in the region immediately centromeric to human MHC across the 6p21.2-6p21.3 chromosomal boundary.					
JOURNAL	Unpublished (1997)					
COMMENT	On Oct 19, 1998 this sequence version replaced gi:2045115.					
	Contact: Tripodis, Nikos					
	Division of Medical and Molecular Genetics					
	Guy's Hospital					
	7th floor, Guy's Tower, London SE1 9RT, UK					
	Email: nikos@nrl.nl					
FEATURES	Location/Qualifiers					
source	1..2275					
	/organism="Homo sapiens"					
	/db_xref="taxon:9606"					
	/map="6p21.3"					
	/clone="ntcon2 contig"					
	/clone_1lb="Human mRNA (Tripodis and Ragoussis)"					
BASE COUNT	438 a 619 c 470 g 599 t 149 others					
ORIGIN						
	Query Match					
	Best local similarity 16.28; Pred. No. 3,49e-10;					
Matches	11; Conservative 39; Mismatches 17; Indels 1; Gaps 1					
	11.3%; Score 27; DB 20; Length 2275;					
Db	1556 WTCMCTSKASASACAMRMKMGMS-RSSRSITWGSMSGCTGTRKRYRYSMTGTRK 1614					
Oy	24 CTCATCCCGAGGAGCGGACACATATGGGCTGCGCGCCCTCTCTCTCTTG 83					
Db	1615 WTTWMMYM 1622					
Oy	84 ACCCGCT 91					
RESULT	8	F06958	348 bp	mRNA	EST	20-FEB-1995
LOCUS	F06958	HSLOC101 normalized infant brain cDNA Homo sapiens	CDNA	clone		
DEFINITION	c-1gc10, mRNA sequence.					
ACCESSION	F06958					
NID	9672595					
VERSION	F06958.1	GI:672595				
KEYWORDS	EST.					
SOURCE	human.					
ORGANISM	human sapiens					
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;					
	Eutheria; Primates; Catarrhini; Hominoidea; Homo.					
REFERENCE	1 (bases 1 to 348)					
AUTHORS	Auffray, C., Benar, G., Bois, F., Bouchier, C., da Silva, C.,					
	Devignes, M.D., Dupret, S., Houligatte, R., Jumeau, M.N., Lamy, B.,					
	Lorenzo, F., Michelli, H., Mariage-Samson, R., Pietu, G., Poullet, Y.,					
	Sebastien-Kabich, C. and Tessier, A.					
TITLE	IMAGE: molecular integration of the analysis of the human genome and its expression					

JOURNAL
MEDLINE
59277534
COMMENT On Sep 21, 1992 this sequence version replaced gi:279286.

Contact: Genethon
Genethon Centre de recherche sur le Genome Humain
1, rue de l'Internationale, Bp60 91002 Evry Cedex, FRANCE
Tel: 33169472800
Fax: 33160778698
Email: genexpress@genethon.fr

Single read
Genexpress_library_id: C; Genexpress_sequence_id: ylc-1q10
Insert Length: 639 Std Error: 0.00
Seq primer: (-21)M13-universal
High quality sequence stop: 150.

FEATURES

SOURCE

1..348
Location/Qualifiers
/organism="Homo sapiens"
/note="Organ: brain; Vector: lafmid BA; Site_1: HindIII;
Site_2: NotI; sex=Female; dev_stage=3 months old;
Isolate=muscular atrophy patient; tissue_type=total
brain; total mRNA was oligo-(dT) primed and directionally
cloned 5' -> 3' into the HindIII -> NotI sites of the
lafmid BA vector. Clone library from B.Soures, psychiatry
Dept. Columbia University, USA. Normalization method:
Bento Soares, P.N.A.S. in press"
/db_xref="taxon:9606"
/clone_id="c-1q10"
/clone_lib="normalized infant brain CDNA"
/sex="Female"
/tissue_type="total brain"
/dev_stage="3 months old"
BASE COUNT 84 a 91 c 90 g 81 t 2 others
ORIGIN

Query Match 10.18; Score 24; DB 19; Length 348;
Best Local Similarity 67.28; Pred. No. 2.31e-06;
Matches 45; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

Db 5 TGTTCCTCCCTCCAGAGATCCCTTGTGTAGTATGCTTACAGTACACACACAC 64
||| ||||| ||||| ||| ||| ||||| ||| ||||| ||| ||||| |||
OY 18 TGTCTCTCCATCCAGAGACGACGACACTATGGGCTGTGGCTCCCTTGTCTCTC 77
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Db 65 CTCTAGA 71
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
OY 78 CTCTTGA 84

RESULT 9
LOCUS AA376266 238 bp mRNA EST 21-APR-1997
DEFINITION EST88915 HSC172 cells II Homo sapiens cDNA 5' end, mRNA sequence.
ACCESSION AA376266
NID 92028809
VERSION AA376266.1 GI:2028809
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 238)

REFERENCE
AUTHORS
Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fullner,R.A.,
Bult,C.J., Lee,N.H., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D.,
White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-Wai,C.,
Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,
Fitzgerald,L.M., Fitzhugh,W.M., Fritchman,J.L., Geoghegan,N.S.,
Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S., Jr.,
Kelley,J.M., Kelley,J.C., Liu,L.-I., Marrairos,S.M., Merrick,J.M.,
Moreno-Palanges,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M.,
Phillips,C.A., Ryder,S.E., Scott,J.L., Saudex,D.M., Shirley,R.,
Small,K.V., Spriggs,T.A., Uterback,T.R., Weidman,J.F., Li,Y.,
Bednarek,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,
Dinhe,D., Feng,D.-F., Fertie,A., Fischer,C., Hastings,G.A.,
He,W.W., Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K.,

TITLE
JOURNAL
MEDLINE
59277534
COMMENT Initial assessment of human gene diversity and expression patterns
based upon 83 million nucleotides of cDNA sequence
Nature 377 (6547 Suppl), 3-174 (1995)
Other ESTs: THC191210
Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA
Tel: 3016699056
Fax: 3016699423
Email: arkerlavage@tigr.org

For clone availability, additional sequence and expression
information related to this EST, please check the TIGR Human Gene
Index (<http://www.tigr.org/tdb/hgi/hgi.html>)
Seq primer: M13 Reverse.

Location/Qualifiers
1..238

/organism="Homo sapiens"
/note="Organ: lung; Vector: pBluescript SK-; Site_1:
EcoRI; Site_2: XhoI"
/db_xref="ATCC (host):180944"
/db_xref="taxon:9606"
/clone_id="HSC172 cells II"
/cell_type="fibroblast"
/cell_line="HSC172 (60PDC)"
/dev_stage="fetal"
BASE COUNT 48 a 70 c 50 g 66 t 4 others
ORIGIN

Query Match 9.7%; Score 23; DB 12; Length 238;
Best Local Similarity 67.2%; Pred. No. 3.76e-05;
Matches 45; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

Db 141 TGTTCCTCCCTCCAGAGATCCCTTGTGTAGTATGCTTACAGTACACACAC 200
||| ||||| ||||| ||| ||| ||||| ||| ||||| ||| ||||| |||
OY 18 TGTCTCTCCATCCAGAGACGACGACACTATGGGCTGTGGCTCCCTTGTCTCTC 77
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Db 201 CTCTAGA 207
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
OY 78 CTCTTGA 84

RESULT 10
LOCUS AA323964 311 bp mRNA EST 20-APR-1997
DEFINITION EST28616 Cerebellum II Homo sapiens cDNA 5' end, mRNA sequence.
ACCESSION AA323964
NID 91976290
VERSION AA323964.1 GI:1976290
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 311)

REFERENCE
AUTHORS
Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fullner,R.A.,
Bult,C.J., Lee,N.H., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D.,
White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-Wai,C.,
Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,
Fitzgerald,L.M., Fitchhugh,W.M., Fritchman,J.L., Geoghegan,N.S.,
Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S., Jr.,
Kelley,J.M., Kelley,J.C., Liu,L.-I., Marrairos,S.M., Merrick,J.M.,
Moreno-Palanges,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M.,
Phillips,C.A., Ryder,S.E., Scott,J.L., Saudex,D.M., Shirley,R.,
Small,K.V., Spriggs,T.A., Uterback,T.R., Weidman,J.F., Li,Y.,
Bednarek,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,
Dinhe,D., Feng,D.-F., Fertie,A., Fischer,C., Hastings,G.A.,
He,W.W., Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K.,

Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA
Tel: 3018699056
Fax: 3018699423
Email: arkerlav@tigr.org
For clone availability, additional sequence and expression
information related to this EST, please check the TIGR Human Gene
Index (<http://www.tigr.org/cdb/hgi/hgi.html>)
Seq primer: M13 Reverse

Query Match	9.7%	Score 23	DB 11	Length 311
Best Local Similarity	89.7%	Pred. No. 3.76e-05		
Matches	26	Conservative	0	Mismatches 3
			Indels	0
			Gaps	0

Contact: Maria M/Mouse EST Project
WASHU-HIMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mousse@watson.wustl.edu
This clone is available royalty-free through LNL ; contact the

```

/Note="Vector: pF73D-pac (pharmacia) with a modified
polyLinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was primed with a Not I -oligo(dT) primer; 15',
TGTTACCAATCTGAAGTGGAGGAGGGCCGGGAATTTTTTTTTTTTTTTTTTT
T 3'], on equal amounts of mRNA from 2 13.5dpc and 2
14.5dpc embryos (total RNA provided by Minoru Ko, Wayne
State Univ., from 2 ); double-stranded cDNA was ligated to
Eco RI adaptors (Pharmacia), digested with Not I and
cloned into the Not I and Eco RI sites of the modified
pF733 vector. Library went through one round of
normalization, and was constructed by Bento Soares and
M. Fatima Bonaldo.
"/
/db_xref="taxon:10090"
/clone IMAGE:352493"
/clone_1lb="Soares mouse embryo NBME13.5 14.5"
/sex="unknown"
/tissue_type="embryo"
/dev_stage="13.5-14.5dpc total fetus"
/lab_host="DH10B"

```

[illegible]

Contact: Robert Strausberg, Ph.D.
 Tel: (301) 496-1550
 Email: Robert.C.Strausberg@nih.gov
 CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
 BonalDO, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CCAP clone distribution Information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bdnp/image/image.html
 Insert Length: 691 Std Error: 0.00
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 402.
 Location/Qualifiers

1. .412

TGTATACATCTGATGAGGAGCGCCGCCCAATTTTTTTTTTTTTTTT 3'].
 Double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified pT733 vector. Library
 went through one round of normalization to cG5, and was
 constructed by Bento Soares and M. Fatima Donaldso.
 db_xref=taxon:9606

BASE COUNT ORIGIN	111 a	78 c	116 g	107 t
----------------------	-------	------	-------	-------

9.78; Score 23; DB 18; Length 412;
Prod No 375205

Matches	45; Conservative	0; Mismatches	22; Indels	0; Gaps	0;
---------	------------------	---------------	------------	---------	----

D6 262 TCATAGAGGTGGTGGTGTCATCCGTAACACCATACTCACCAGGAATCGGGAGGCA 3
|| ||||| || || ||||| || || ||||| ||
CP 84 TCAAAGGAGGACAAAGGGGCAGCCGACGCCCATAGTGTGGCCACTGCTCTCTGGGATGGA 29

Db	322	GGCAACA	32
Cp	24	GGAGACA	18

RESULT	13	
LOCUS	T15255	426 bp mRNA
DEFINITION	cr855 lambdaZAPR	EST
	sequence.	28-JUL-1995
		Ricinus communis cDNA clone pcr855, mRNA

VERSION T15255.1 GI:688907
KEYWORDS EST.

ORGANISM

Euairycal; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta
euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;
Rosidae; Euphorbiales; Euphorbiaceae; Ricinus.
1 (base) + 0.435.

AUTHORS	TITLE	JOURNAL	COMMENT
vandeloo, F.J., Turner, S. and Somerville, C.	Expressed sequence tags from developing castor seeds	Plant Physiol. 108, 1141-1150 (1995)	

Contact: Somerville CR
Carnegie Institution
Carnegie Institution, 290 Panama St., Stanford, CA 94305
Tel: 4153251521
Email: ccsdandrew.stanford.edu
seq primer: T3.

FEATURES

Location/Qualifiers
1. .426

```
/organism="Ricinus communis"
/strain="Baker 296"
```

/not=Vector; lambdaZAP11; Site 1: EcoRI, Site 2: XhoI;
 Poly(A)+ RNA was purified from developing stage III to
 stage V (Greenwood & Bevely, Can. J. Bot. 60:1151-1160,
 1982) endosperm plus embryo of immature cotton fruits.
 cDNA was synthesized and cloned into lambdaZAP11 according
 to the instructions of the manufacturer (Stratagene);
 synthesis was primed from the Poly(A) tail, and cloned
 directionally into XhoI (3') and EcoRI (5') sites. In few
 cases, sequence data indicated that this directionality
 was reversed. Partial cDNA clones predominate."
 db_xref="tacon:3986"

BASE COUNT	126 a	61 c	99 g	129 t	11 others
ORIGIN	/clone="pcrs855" /clone_lib="lambdaZAPST"				

Query Match	9.7%;	Score 23;	DB 8;	Length 426;
Best Local Similarity	71.4%;	Pred. No. 3.76e-05;		
Matches	35;	Conservative	0;	Mismatches 14;
			Indels	0;
			Gaps	0

Db 77 TGACCATTCACAGTACCTGAAGATGACTGTCTGANTGATTTGNCCTA 122
 |||| | ||| ||||| |||| | |||| ||| ||||
 Qy 121 TGAATTGCACACTGAACTGAGGAGCTTTTTCGACAAATTCCTCTA 166

RESULT	14				
LOCUS	AA498488	481 bp	MRNA	EST	01-JUL-1997
DEFINITION	vha4b06.r1 Barstead mouse pooled organs MFLRB4 Mus musculus c				

clone IMAGE:889811 5' similar to gb:M30514 mouse muscle nicotinic acetylcholine receptor gamma-subunit (mouse);, mRNA sequence

```

NID          92233511
VERSION      AA498488.1  GI:2233511

```

KEYWORDS	EST.
SOURCE	house mouse.

ORGANISM Mus musculus

REFERENCE
1 (bases 1 to 481)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS

Geisels, S., Kincaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schei, I., Schellberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Treising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and Meterson, R.

TITLE	ine wasnu-HHMI Mouse EST Project
JOURNAL	Unpublished (1996)
COMMENT	On Sep 12, 1996 this sequence version replaced gi.1405167

Contact: Maria M/Mouse EST Project
MashU-HHMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.edu

This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
MGI:517771

Seq primer: -28ml3 rev2 ET from Amersham
High quality sequence stop: 461.

FEATURES	Location/Qualifiers
source	1. .481

1.481

/organism="Mus musculus"

```
/note="Organ: pooled; Ve
```

strand cDNA was primed w

3']; double-stranded CD

The Not I and Eco RI site

```
/db_xref="taxon:10090"
```

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/clone_lib="Barstead mou
```

```
/tissue_type="pooled org
```

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/1ab_nosf="DH10B"
131 ~ 134 ~ 10

```

BASE COUNT	120 a	131 c	124 g	106 t
ORIGIN				

```
Query Match      9.7%; Score 23; DB 14; Length 481
Best Local Similarity 76.7%; Pred. No. 3.76e-05;
```

Matches 33: Conservative 0: Mismatches 10: Indels 0: Gaps 0:

Db 277 CTTCAAGTGTCAACACCTGCTGAGCCCGACGTCATATGG 319
 12 CTTCAAGTGTCTCCATCCAGAGCGCGACGTCATATGG 54

RESULT 15

LOCUS AF038250 1287 bp mRNA EST 30-MAR-1998
 DEFINITION AF038250 Human mRNA (Tripodis and Ragousis) Homo sapiens CDNA
 clone ntcon9, mRNA sequence.

ACCESSION AF038250

NID 92815880

VERSION AF038250.1 GI:2815880

KEYWORDS EST

SOURCE human.

ORGANISM

Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE

1 (bases 1 to 1287)
 Tripodis, N. and Ragousis, J.

Generation of a transcription map in the region immediately
 centromeric to human MHC across the 6p21.2-6p21.3 chromosomal
 boundary

Unpublished (1997)

JOURNAL

COMMENT On Jan 19, 1998 this sequence version replaced gi:2045085.

Contact: Tripodis, Nikos
 Division of Medical and Molecular Genetics
 Guys Hospital
 7th floor, Guy's Tower, London SE1 9RT, UK
 Email: nikosenk1.n1.

FEATURES

source

location/Qualifiers
 1..1287
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /map="6p21.3"
 /clone="ntcon9"
 /clone_1b="Human mRNA (Tripodis and Ragousis)"

BASE COUNT 349 a 219 c 293 g 361 t 65 others

ORIGIN

Query Match

9.7%; Score 23; DB 20; Length 1287;

Best Local Similarity 25.8%; Pred. No. 3,76e-05;

Matches 24: Conservative 32: Mismatches 37: Indels 0: Gaps 0:

Db 381 VVATCAGSCCVCACCTGBCDCTGGGTBSHBVCMCBANADGATBACGKGVGA 440

53 GGAGTCTGGGGTGGCCCTGCTCTCTCTGACCTCTGCGACCTCAGATGAGACAGG 112

Db 441 STCTHNCDDCKTGGSGAGTYVNHDMNMGGA 473

113 GCCGGATGACTTGTGCACTGAAGCTGAAGGA 145

Search completed: Sun Oct 24 16:38:34 1999
 Job time : 471 secs.

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L27 ANSWER 1 OF 4 CAPLUS COPYRIGHT 1999 ACS DUPLICATE 1
ACCESSION NUMBER: 1999:518247 CAPLUS
DOCUMENT NUMBER: 131:166197
TITLE: Methods for detecting lung
diseases
INVENTOR(S): Cohen, Maurice; Friedman, Paula
N.; Gordon, Julian; Hodges, Steven C.;
Klass, Michael R.; Kratochvil, Jon D.;
Roberts-Rapp, Lisa; Russell, John C.;
Stroupe, Steven D.
PATENT ASSIGNEE(S): Abbott Laboratories, USA
SOURCE: U.S., 36 pp., Cont.-in-part of U.S. Ser. No.
744,211, abandoned.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5939265	A	19990817	US 1997-964725	19971105
PRIORITY APPLN. INFO.:			US 1996-744211	19961105

AB A set of contiguous and partially overlapping RNA sequences and polypeptides encoded thereby, designated as LU103 and transcribed from lung tissue are described. A fully sequenced clone representing the longest continuous sequence of LU103 is also disclosed. These sequences are useful in detecting, diagnosing, staging, monitoring, prognosticating, preventing or treating, or detg. the predisposition of an individual to diseases and conditions of the lung such as lung cancer.

L27 ANSWER 2 OF 4 CAPLUS COPYRIGHT 1999 ACS DUPLICATE 2
ACCESSION NUMBER: 1999:8152 CAPLUS
DOCUMENT NUMBER: 130:77055
TITLE: Protein LS170 and cDNA sequences useful for
detecting diseases of the human
lung
INVENTOR(S): Billing-Medel, Patricia A.;
Cohen, Maurice; Colpitts, Tracey
L.; Friedman, Paula N.; Gordon,
Julian; Granados, Edward N.; Hodges, Steven C.;
Searcher : Shears 308-4994

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Klass, Michael R.; Kratochvil, Jon D.;
Roberts-Rapp, Lisa; Russell, John C.;
Stroupe, Stephen D.

PATENT ASSIGNEE(S): Abbott Laboratories, USA
SOURCE: PCT Int. Appl., 120 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9856951	A1	19981217	WO 1998-US11601	19980611
W: CA, JP				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				

PRIORITY APPLN. INFO.: US 1997-49183 19970611

AB A set of contiguous and partially overlapping cDNA sequences and polypeptides encoded thereby, designated as LS170 and transcribed from human lung tissue, is described. These sequences are useful for the detecting, diagnosing, staging, monitoring, prognosticating, in vivo imaging, preventing or treating, or detg. the predisposition of an individual to diseases and conditions of the lung, such as lung cancer. Also provided are antibodies which specifically bind to a LS170-encoded polypeptide or protein, and agonists or inhibitors which prevent action of tissue-specific LS170 polypeptides, which mols. are useful for the therapeutic treatment of lung diseases, tumors, or metastases.

L27 ANSWER 3 OF 4 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD

ACCESSION NUMBER: 1998-437479 [37] WPIDS

DOC. NO. NON-CPI: N1998-340776

DOC. NO. CPI: C1998-133112

TITLE: New nucleic acid for the lung
disease marker LU105 - polypeptides,
antibodies and genes, used for diagnosis,
prevention, treatment of lung
disease, specifically cancer.

DERWENT CLASS: B04 D16 S03

INVENTOR(S): BILLING-MEDEL, P A; COHEN, M;
COLPITTS, T L; FRIEDMAN, P N;
GORDON, J; GRANADOS, E N; HODGES, S C; KLASS,
M R; KRATOCHVIL, J D; ROBERTSRAPP, L;
RUSSELL, J C; STROUPE, S D

PATENT ASSIGNEE(S): (ABBO) ABBOTT LAB

COUNTRY COUNT: 19

PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9833926	A1	19980806	(199837)*	EN	117
RW: AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE					
W: CA JP					

APPLICATION DETAILS:

Searcher : Shears 308-4994

PATENT NO	KIND	APPLICATION	DATE
WO 9833926	A1	WO 1998-US1766	19980130

PRIORITY APPLN. INFO: US 1997-791710 19970131

AN 1998-437479 [37] WPIDS

AB WO 9833926 A UPAB: 19980916

A method for detecting target LU105 nucleic acid (I) comprises treating a sample with at least one LU105-specific nucleic acid (II), or its complement. (II) is at least 50% identical with 190, 244, 225, 114, 562 or 519 bp sequences given in the specification, or their fragments and complements.

Also claimed are:

(1) (I) or its fragments able to hybridise selectively to the LU105 gene and having at least 50% identity with the 190, 244, 225, 114, 562 or 519 bp sequences given above;

(2) recombinant expression systems including (I) and control sequence;

(3) cells transformed with this expression system;

(4) LU105 polypeptides (III) at least 50% identical with the 104, 26, 19, 21, 18 or 19 amino acid (aa) sequences given in the specification or their fragments;

(5) antibodies (Ab) that bind to at least one LU105 epitope present in (III);

(6) cells transformed with the 190, 244, 225, 114, 562 or 519 bp sequences described above;

(7) LU105 specific nucleic acid (II); and

(8) genes, or their fragments, that encode a protein at least 50% identical with the 104 aa sequences as in (4).

USE - LU105 is a **lung disease** marker. Cells as in (3) are used to express recombinant (III) which are used to raise Ab. Ab are used to detect the LU105 antigen, and correspondingly this antigen is used to detect specific antibodies, in usual immunoassays. (I) and (III) are used for diagnosis, staging, monitoring, prognosis, prevention, treatment (e.g. using antisense molecules, ribozymes, Ab or other antagonists) and determination of susceptibility to, **lung disease**, specifically cancer. (III) are also used to screen for specific binding agents, potentially useful therapeutically. LU105 is a marker for **lung disease** (present at high concentration, in altered form or in an unusual body compartment).

ADVANTAGE - LU105 can be detected in blood, plasma or serum in an inexpensive, non-invasive test.

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L27 ANSWER 4 OF 4 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD

ACCESSION NUMBER: 1998-286957 [25] WPIDS

DOC. NO. NON-CPI: N1998-225472

DOC. NO. CPI: C1998-088988

TITLE: Lung tissue derived polynucleotide LU103 - useful to detect, diagnose, stage, monitor, prognose, prevent, treat or determine pre-disposition to **lung disease**, e.g. **lung cancer**.

Searcher : Shears 308-4994

DERWENT CLASS: B04 D16 S03
 INVENTOR(S): COHEN, M; FRIEDMAN, P N;
 GORDON, J; HODGES, S C; KLASS, M R;
 KRATOCHVILL, J D; ROBERTS-RAPP, L; RUSSELL, J
 C; STROUPE, S D
 PATENT ASSIGNEE(S): (ABBO) ABBOTT LAB
 COUNTRY COUNT: 19
 PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9820143	A1	19980514	(199825)*	EN	86
RW: AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE					
W: CA JP					

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9820143	A1	WO 1997-US20680	19971105

PRIORITY APPLN. INFO: US 1996-744211 19961105

AN 1998-286957 [25] WPIDS

AB WO 9820143 A UPAB: 19980715

The following are claimed: (1) a method for detecting the presence of a target LU103 polynucleotide in a test sample, comprising: (a) contacting the sample with at least 1 LU103-specific polynucleotide, and (b) detecting the target LU103 polynucleotide in the test sample, where the LU103 polynucleotide has at least 50% identity to the 269, 263, 225, 507 or 519 bp nucleic acid sequence given in the specification; (2) a method for detecting LU103 mRNA in a test sample, comprising: (a) performing reverse transcription with at least 1 primer in order to produce cDNA; (b) amplifying the cDNA using LU103 oligonucleotides as sense and antisense primers to obtain LU103 amplicon, and (c) detecting LU103 amplicon in the test sample, where the LU103 oligonucleotides utilised in steps (a) and (b) have at least 50% sequence identity to the 269, 263, 225, 507 or 519 bp sequence and (3) a method detecting a target LU103 polynucleotide in a test sample suspected of containing the target, comprising: (a) contacting the test sample with at least 1 LU103 oligonucleotide as a sense primer and at least 1 LU103 oligonucleotide as an anti-sense primer and amplifying to obtain a first stage reaction product; (b) contacting the first stage reaction product with at least 1 other LU103 oligonucleotide to obtain a second stage reaction product, provided that the other LU103 oligonucleotide is located 3' to the LU103 oligonucleotides utilised in step (a) and is complementary to the first stage reaction product, and (c) detecting the second stage reaction product as an indication of the presence of the target LU103 polynucleotide, where the LU103 oligonucleotides utilised in steps (a) and (b) have at least 50% sequence identity to the 269, 263, 225, 507 or 519 bp sequence; (4) a purified polynucleotide derived from an LU103 gene, where the polynucleotide is capable of selectively hybridising to the nucleic acid of the LU103 gene and has at least 50% identity to the 269, 263, 225, 507 or 519 bp

Searcher : Shears 308-4994

- sequence; (5) a recombinant expression system comprising a nucleic acid sequence that includes an ORF derived from LU103 operably linked to a control sequence compatible with a desired host, where the nucleic acid sequence has at least 50% identity to the 269, 263, 225, 507 or 519 bp sequence; (6) cell transfected with the recombinant expression system; (7) cell transfected with a nucleic acid sequence encoding at least 1 LU103 epitope, where the nucleic acid sequence has the 269, 263, 225, 507 or 519 bp sequence; (8) composition comprising a LU103 polynucleotide, where the polynucleotide has at least 50% identity to the 269, 263, 225, 507 or 519 bp sequence; (9) gene which encodes a LU103 protein which comprises an amino acid sequence with at least 50% identity with the 93 residue amino acid sequence given in the specification, and (10) gene comprising DNA having at least 50% identity with the 507 or 519 bp sequence.

USE - The methods and products of the invention may be used to detect, diagnose, stage, monitor, prognose, prevent, treat or determine the predisposition diseases and conditions of the lung, e.g. lung cancer.

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(FILE 'CAPLUS' ENTERED AT 12:09:17 ON 25 OCT 1999)

L1 10217 SEA ABB=ON PLU=ON LUNG(3A)(DISEAS? OR DISORDER)
L2 583 SEA ABB=ON PLU=ON L1(S)(IDENTIF? OR DETECT? OR DET##
OR DETERM? OR DIAGNOS?)
L3 14 SEA ABB=ON PLU=ON L2 AND (REAGENT OR EPITOPE OR LS147
OR LS 147)

L3 ANSWER 1 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1999:556186 CAPLUS

TITLE: Pigeon fanciers' lung:
identification of disease

-associated carbohydrate epitopes on
pigeon intestinal mucin

AUTHOR(S): Baldwin, C. I.; Todd, A.; Bourke, S. J.; Allen,
A.; Calvert, J. E.

CORPORATE SOURCE: Department of Immunology, The Medical School,
University of Newcastle upon Tyne, Newcastle
upon Tyne, NE2 4HH, UK

SOURCE: Clin. Exp. Immunol. (1999), 117(2), 230-236
CODEN: CEXIAL; ISSN: 0009-9104

PUBLISHER: Blackwell Science Ltd.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Pigeon intestinal mucin, a complex high mol. wt. glycoprotein, is a key antigen in the development of pigeon fanciers' lung (PFL). We have studied the specificity of antibodies to mucin in patients with PFL and asymptomatic antibody-pos. individuals. Extensive papain digestion, which removes the non-glycosylated regions of the mucin leaving the heavily glycosylated "bottle brush" regions, resulted in a 600-fold decrease in IgG3 antibody titers with little effect on IgG1 and IgG2 titers. This suggests that IgG1 and IgG2 are directed against the region rich in O-linked sugar chains while the majority of the IgG3 is directed against **epitopes** which are proteinase-sensitive. Lectin mapping of the carbohydrates present on pigeon intestinal mucin demonstrated high levels of exposed N-acetyl neuraminic acid, N-acetyl galactosamine and N-acetyl glucosamine, with lower levels of fucose and some galactose. Sera from pigeon fanciers inhibited binding of lectins specific for N-acetyl neuraminic acid, N-acetyl galactosamine, internal N-acetyl glucosamine and fucose. Sera from people with PFL, compared with sera from asymptomatic antibody-pos. fanciers, had significantly higher titers of antibody that inhibited binding of four lectins specific for N-acetyl galactosamine and one fucose-specific lectin, suggesting that these sugars may play a dominant role in disease-assocd. **epitopes**. The results suggest that different IgG subclasses recognize different **epitopes** on mucin and that the **epitopes** recognized by the major subclasses are present on the O-linked oligosaccharides. Further, the carbohydrate-specific anti-mucin antibodies produced by PFL

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patients may differ in their specificity from those found in asymptomatic individuals.

L3 ANSWER 2 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1999:8152 CAPLUS

DOCUMENT NUMBER: 130:77055

TITLE: Protein LS170 and cDNA sequences useful for
detecting diseases of the
human lung

INVENTOR(S): Billing-Medel, Patricia A.; Cohen, Maurice;
Colpitts, Tracey L.; Friedman, Paula N.; Gordon,
Julian; Granados, Edward N.; Hodges, Steven C.;
Klass, Michael R.; Kratochvil, Jon D.;
Roberts-Rapp, Lisa; Russell, John C.; Stroupe,
Stephen D.

PATENT ASSIGNEE(S): Abbott Laboratories, USA

SOURCE: PCT Int. Appl., 120 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9856951	A1	19981217	WO 1998-US11601	19980611
W: CA, JP				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				

PRIORITY APPLN. INFO.: US 1997-49183 19970611

AB A set of contiguous and partially overlapping cDNA sequences and polypeptides encoded thereby, designated as LS170 and transcribed from human lung tissue, is described. These sequences are useful for the detecting, diagnosing, staging, monitoring, prognosticating, in vivo imaging, preventing or treating, or detg. the predisposition of an individual to diseases and conditions of the lung, such as lung cancer. Also provided are antibodies which specifically bind to a LS170-encoded polypeptide or protein, and agonists or inhibitors which prevent action of tissue-specific LS170 polypeptides, which mols. are useful for the therapeutic treatment of lung diseases, tumors, or metastases.

L3 ANSWER 3 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1998:674032 CAPLUS

DOCUMENT NUMBER: 130:64487

TITLE: Immunohistochemical detection of multidrug
resistance protein in human lung cancer and
normal lung

AUTHOR(S): Wright, Scott R.; Boag, Alexander H.;
Searcher : Shears 308-4994

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Valdimarsson, Gunnar; Hipfner, David R.;
Campbell, Barbara G.; Cole, Susan P. C.; Deeley,
Roger G.
CORPORATE SOURCE: Departments of Pathology and Cancer Research
Laboratories, Queen's University, Kingston, ON,
K7L 3N6, Can.
SOURCE: Clin. Cancer Res. (1998), 4(9), 2279-2289
CODEN: CCREP4; ISSN: 1078-0432
PUBLISHER: American Association for Cancer Research
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Monoclonal antibody QCRL-1 is highly specific for a defined linear
epitope in a relatively poorly conserved region of the human
multidrug resistance protein (MRP). The authors have used QCRL-1 to
examine MRP expression in archival and fresh snap-frozen samples of
untreated small cell (SC) and non-small cell (NSC) lung cancers
(LCs), as well as normal lung. The authors found that the majority
(87%) of all histol. subtypes of NSCLC had detectable levels of MRP
in most of the tumor mass. In a substantial proportion of
adenocarcinomas (55%) and squamous cell carcinomas (28%),
immunoreactivity approached that obtained with the highly multidrug
resistant cell line H69AR from which the MRP was originally cloned.
Both the level and frequency of MRP expression in untreated SCLC was
significantly lower than in NSCLC. The MRP was detectable in only
56% of SCLC tumors and, in most cases, was expressed in small focal
clusters of cells. Immunofluorescence studies of tumor tissue and
normal lung confirmed the plasma membrane location of the MRP.
However, in normal bronchial epithelium and seromucous glands,
unlike in tumor cells, the MRP was detected only on basolateral
membranes. In addn., strong MRP immunoreactivity was detected in
reactive type II pneumocytes present in hyperplastic alveoli, but
not in normal type I and type II pneumocytes. No potentially
confounding correlation independent of its possible role in drug
resistance was obsd. between MRP expression in untreated NSCLC and
any clinicopathol. parameter examd., including overall survival.

L3 ANSWER 4 OF 14 CAPLUS COPYRIGHT 1999 ACS
ACCESSION NUMBER: 1997:692333 CAPLUS
DOCUMENT NUMBER: 127:316564
TITLE: Diagnosis of chronic respiratory failure and
pulmonary emphysema, and diagnostic
reagents
INVENTOR(S): Okuda, Yukichi; Morita, Riichiro; Hanatani,
Mitsuya; Matsuo, Katsuhiko
PATENT ASSIGNEE(S): Toa Gosei Chemical Industry Co., Ltd., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 4 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
Searcher : Shears 308-4994

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FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	JP 09274037	A2	19971021	JP 1996-104629	19960402
AB	Detn. of human serum vascular endothelial growth factor/vascular permeability factor (VPF) is useful for accurate diagnosis of the title diseases. The detn. is carried out by using anti-VPF antibodies or VPF receptors. The serum level of VPF in patients with the diseases is much higher than healthy humans.				

L3 ANSWER 5 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1997:44677 CAPLUS

DOCUMENT NUMBER: 126:55930

TITLE: Detection of neoplastic cells based on alternatively spliced transcripts of the p15INK4B and p16INK4A cyclin/CDK inhibitors
INVENTOR(S): Sidransky, David; Baylin, Stephen B.
PATENT ASSIGNEE(S): Johns Hopkins University School of Medicine, USA
SOURCE: PCT Int. Appl., 82 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	WO 9635704	A1	19961114	WO 1996-US6666	19960510
	W: AU, CA, CN, JP, KR RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	US 5767258	A	19980616	US 1995-439962	19950512
	US 5856094	A	19990105	US 1995-497535	19950630
	AU 9657399	A1	19961129	AU 1996-57399	19960510
PRIORITY APPLN. INFO.:				US 1995-439962	19950512
				US 1995-497535	19950630
				WO 1996-US6666	19960510
AB	Novel cell cycle regulatory polynucleotide transcripts and their encoded polypeptides are provided which were identified as the products of alternatively spliced mRNA for cyclin/CDK inhibitors, p16INK4A and p15INK4B, and a 5' nucleotide sequence referred to as 5'ALT. The p16 and p15 genes colocalize to human chromosome 9p21, which has been identified as a region having homozygous deletions in many tumors. 5'ALT also resides on chromosome 9p21, just 5' of exon 2 of p15, and about 30 kb upstream from p16. Polynucleotide transcripts are provided in which a 5'ALT polynucleotide is operatively linked to (1) exon 2 and exon 3 of p15 or (2) exon 2 of				

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p15. These transcripts are assocd. with normal growth control and regulation of cellular proliferation and provide a means for the development of more accurate diagnostic, prognostic, and therapeutic regimes for disorders assocd. with control of cell cycle progression and cell differentiation and the loss of such control. Methylation of p16 DNA and a resultant decrease in p16 gene expression is assocd. with transcriptional block and is assocd. with a variety of neoplasms. Thus, a method for detecting a neoplasm in a subject by detecting methylation of 5'CpG islands in p16 DNA, or detecting p16 mRNA or polypeptide levels in a sample is also provided. Preferably, the method utilizes a methylation-sensitive restriction endonuclease in order to detect p16 methylation. The 5'ALT, or 5'ALT-p16 or 5'ALT-p15 polypeptides can also be used to produce antibodies which are immunoreactive or bind to **epitopes** of the resp. polypeptides.

L3 ANSWER 6 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1996:68700 CAPLUS

DOCUMENT NUMBER: 124:169878

TITLE: Difference in sero-diagnostic values among
KL-6-associated mucins classified as Cluster 9

AUTHOR(S): Kohno, Nobuoki; Inoue, Yoshikazu; Hamada,
Hironobu; Fujioka, Seiji; Fujino, Shun;
Yokoyama, Akihito; Hiwada, Kunio; Ueda,
Norifumi; Akiyama, Mitoshi

CORPORATE SOURCE: School Medicine, Ehime University, Ehime,
791-02, Japan

SOURCE: Int. J. Cancer, Suppl. (1994), 8(Third
International IASLC Workshop on Lung Tumor and
Differentiation Antigens, 1993), 81-3
CODEN: IJSUEZ; ISSN: 0898-6924

DOCUMENT TYPE: Journal

LANGUAGE: English

AB KL-6 classified as Cluster 9 (MUC-1) is a circulating high-mol.-wt. mucin-like mol. Serum level of KL-6 was measured by a sandwich assay using KL-6 antibody as not only a catcher but also as a tracer. The authors established 2 addnl. monoclonal antibodies (MAbs), LISA 101 and EH-123, reacting with KL-6 **epitopes** different from the epitope recognized by KL-6 antibody. The KL-6-assocd. mucins detected by the sandwich assay using LISA 101 or EH-123 antibody as a catcher and KL-6 antibody as a tracer were designated as LISA 1-6 and CAM 123-6 resp. The diagnostic values as the serum markers of KL-6, LISA 1-6 and CAM 123-6 were evaluated measuring their levels in the same serum from healthy individuals and from patients with pulmonary, pancreatic and breast adenocarcinomas. KL-6 was increased abnormally at high rates of more than 50% in pancreatic cancer and in benign lung diseases, LISA 1-6 only in pancreatic cancer, and CAM 123-6 only in pulmonary adenocarcinoma. In benign lung diseases, however, LISA 1-6 and CAM

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123-6 were increased abnormally at the rates of only 5.3% and 0% resp. These observations clearly indicate that LISA 1-6 and CAM 123-6 constitute a part of KL-6, but that they are superior to KL-6 as tumor markers for pancreatic cancer and for pulmonary adenocarcinoma resp., because of their much lower false-pos. rates.

L3 ANSWER 7 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1994:104481 CAPLUS
DOCUMENT NUMBER: 120:104481
TITLE: Monoclonal antibodies against farmer's lung antigens having specific binding to IgG antibodies
AUTHOR(S): Kumar, Anoop; Elms, Nancy; Kurup, Viswanath P.
CORPORATE SOURCE: Dep. Med., Med. Coll. Wisconsin, V., Milwaukee, WI, USA
SOURCE: Int. Arch. Allergy Immunol. (1993), 102(1), 67-71
CODEN: IAAIEG; ISSN: 1018-2438
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Hypersensitivity pneumonitis resulting from environmental exposure to *Saccharopolyspora rectivirgula* (*Micropolyspora faeni*) among farmers has been well recognized. The diagnosis of the disease depends on demonstration of circulating antibodies against *S. rectivirgula*. However, dependable pure antigens are not available for serodiagnosis. The authors employed hybridoma technol. to obtain monoclonal antibodies against *S. rectivirgula* antigens. These monoclonal antibodies were employed to purify antigens through affinity chromatog. When tested in ELISA, high levels of antibodies were demonstrated against these antigens in farmer's lung patient sera compared to exposed but asymptomatic individuals from the same household. In Western blots, patient sera reacted with components of crude antigens with mol. masses of 28, 35, 60, 65 and 68 kD and 4 components above 100 kD, while the monoclonal antibodies reacted only with the 60-kD protein. These purified antigens can be used as reliable **reagents** in the specific **diagnosis** of farmer's lung diseases.

L3 ANSWER 8 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1993:467319 CAPLUS
DOCUMENT NUMBER: 119:67319
TITLE: Test **reagent** for hemoptysic sputum for diagnosis of lung cancer
INVENTOR(S): Qin, Dexing
PATENT ASSIGNEE(S): Chinese Academy of Medical Sciences, Tumour Hospital, Peop. Rep. China
SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu, 4 pp.
CODEN: CNXXEV
Searcher : Shears 308-4994

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DOCUMENT TYPE: Patent
LANGUAGE: Chinese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
	CN 1069341	A	19930224	CN 1991-105479	19910814
AB	The title reagent contg. alc., H ₂ O ₂ , glacial AcOH, and guaiac gum is used for detecting hidden blood in sputum for alerting a possible lung cancer disease . Using the test reagent , the incidence of lung cancer was 40 times higher in pos.-responding people than in neg.-responding people.				

L3 ANSWER 9 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1993:444470 CAPLUS
DOCUMENT NUMBER: 119:44470
TITLE: Automated determination of crosslinked fibrin derivatives in plasma
AUTHOR(S): Elms, M. J.; Bundesen, P. G.; Rowbury, D.; Goodall, S.; Wakeham, N.; Rowell, J. A.; Hillyard, C. J.; Rylatt, D. B.
CORPORATE SOURCE: Pathol. Dep., R. Brisbane Hosp., Brisbane, Australia
SOURCE: Blood Coagulation Fibrinolysis (1993), 4(1), 159-64
CODEN: BLFIE7; ISSN: 0957-5235
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Automated assays for the measurement of crosslinked fibrin derivs. in plasma (XL-FbDP) were developed by using latex beads coated with anti-D-dimer monoclonal antibody (DD-3B6/22) for both the Cobas Fara Chem. Centrifugal and the Cobas Mira analyzers (Roche, Basle, Switzerland). The analyzers were programmed to mix plasma and latex **reagent** simultaneously and analyze absorbance changes over a 10-15 min period. Results were interpolated by the analyzer from a std. curve derived from a polymer of D-dimer. Both assays had high precision (<5% CV) for values between 100 and 1000 ng/mL and provided clear discrimination between normal samples and samples from patients suffering from the thrombotic diseases deep vein thrombosis/pulmonary embolism and disseminated intravascular coagulation. The results obtained for XL-FbDP detn. with both methods compared well with established methods: a high correlation was obtained with a semiquant. manual latex method for both the Fara (r = 0.92) and Mira (r = 0.83) and correlations (r) of 0.81 (Fara) and 0.84 (Mira) were obtained with an EIA. Correlation between the 2 automated procedures was high (r = 0.96). The automated method will enable labs. to provide a rapid and accurate quantitation of

Searcher : Shears 308-4994

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XL-FbDP.

L3 ANSWER 10 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1992:608482 CAPLUS

DOCUMENT NUMBER: 117:208482

TITLE: Detection of early platelet activation and
prediagnosis of thrombotic events by immunoassay
for platelet surface thrombospondin (TSP)

INVENTOR(S): Aiken, Martha L.; Painter, Richard G.

PATENT ASSIGNEE(S): University of Texas System, USA

SOURCE: PCT Int. Appl., 105 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9215886	A1	19920917	WO 1992-US1757	19920309
W: AT, AU, BB, BG, BR, CA, CH, CS, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, PL, RO, RU, SD, SE				
RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FR, GA, GB, GN, GR, IT, LU, MC, ML, MR, NL, SE, SN, TD, TG				
US 5256538	A	19931026	US 1991-668272	19910308
AU 9216554	A1	19921006	AU 1992-16554	19920309
PRIORITY APPLN. INFO.:			US 1991-668272	19910308
			WO 1992-US1757	19920309

AB Persons at risk for a thrombotic event are identified by early immunol. detn. of elevated platelet surface TSP using a (labeled) monoclonal antibody to TSP. Thus, IgG-coated magnetic beads were exposed to an anti-TSP monoclonal antibody and then mixed with paraformaldehyde-fixed human platelets. After magnetic sepn. of the beads, the no. of platelets remaining in suspension was inversely related to TSP surface expression on the platelets. Diagnostic kits contg. **reagents** for the assay are described.

L3 ANSWER 11 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1992:233421 CAPLUS

DOCUMENT NUMBER: 116:233421

TITLE: Human surfactant protein-A contains blood group
A antigenic determinants

AUTHOR(S): Stahlman, Mildred T.; Gray, Mary E.; Ross, Gary
F.; Hull, William M.; Wikenheiser, Kathryn;
Dingle, Sharon; Zelenski-Low, Kay R.; Whitsett,
Jeffrey A.

CORPORATE SOURCE: Sch. Med., Vanderbilt Univ., Nashville, TN,
37232-2370, USA

SOURCE: Pediatr. Res. (1992), 31(4, Pt. 1), 364-71

Searcher : Shears 308-4994



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CODEN: PEREBL; ISSN: 0031-3998

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A major blood group antigenic **epitope** was identified on human pulmonary surfactant protein A (SP-A). Monoclonal (MAb) and polyclonal antibodies generated against purified human SP-A aggregated blood group A human erythrocytes and immunostained epithelial cells in a variety of human tissues, consistent with the tissue distribution of major blood group antigens. SP-A MAb (MAb-8) agglutinated red cells and immunostained tissues from A or AB blood groups, but did not react with cells or tissues from O or B individuals. MAb-8 immunostaining of tissue from blood group A individuals was ablated by incubation with blood group A red cells. MAb and polyclonal antibodies directed against A blood group antigens reacted strongly with purified SP-A obtained from a blood group A individual with alveolar proteinosis. MAb and polyclonal antibodies specific for B blood group antigen failed to react with SP-A from this patient or from patients who were in blood group B. Reactivity of anti-blood group MAb was lost after treatment of SP-A with endoglycosidase-F, demonstrating its reactivity with an **epitope** dependent on the asparagine-linked oligosaccharide at asparagine 187. Reactivity of MAb-8 with SP-A persisted after endoglycosidase-F treatment, but was lost after digestion with collagenase as assessed by Western blot after SDS-PAGE. Reactivity of MAb to SP-A was sensitive to β -elimination, supporting the presence of another blood group antigenic site distinct from the **epitope** dependent on the asparagine-linked carbohydrate. The finding that the SP-A mol. contains a major blood group **epitope** has implication for the clin. use of surfactant replacement preps. and diagnostic **reagents** based on this protein.

L3 ANSWER 12 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1989:210563 CAPLUS

DOCUMENT NUMBER: 110:210563

TITLE: Monoclonal antibodies to angiotensin-converting enzyme: a powerful tool for lung and vessel studies

AUTHOR(S): Danilov, S.; Sakharov, I.; Martynov, A.; Faerman, A.; Muzykantov, V.; Klibanov, A.; Trakht, I.

CORPORATE SOURCE: Inst. Exp. Cardiol., Cardiol. Res. Cent., Moscow, 121552, USSR

SOURCE: J. Mol. Cell. Cardiol. (1989), 21(Suppl. 1), 165-70

CODEN: JMCDAY; ISSN: 0022-2828

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A series (12 clones) of hybridomas were obtained, which produce
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monoclonal antibodies (Mab) to 5 different epitopes of the angiotensin-converting enzyme (ACE) mol. These antibodies may be used to (1) map antigenic structure of ACE, including the study of immunol. heterogeneity of ACE from different organs and tissues; (2) study the immunohistochem. distribution of ACE in human tissues, including the diagnosis of sarcoidosis; (3) develop an ACE immunoassay, and (4) prep. an immunosorbent for large-scale ACE isolation and for ACE-apheresis. One of the antibodies, 9B9, when injected into the circulation of rat and monkey, accumulated with high specificity in the lungs as compared with either normal mouse IgG or other organs and blood. The highly specific and nontoxic accumulation of Mab 9B9 suggests that it also may be used for .gamma. scintigraphy visualization of the pulmonary vascular bed, detection of lung injury, and as a vector for targeted drug delivery to the lung.

L3 ANSWER 13 OF 14 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1988:585013 CAPLUS

DOCUMENT NUMBER: 109:185013

TITLE: Diisocyanate antigens that detect specific antibodies in exposed workers and guinea pigs

AUTHOR(S): Jin, Ruzhi; Karol, Meryl H.

CORPORATE SOURCE: Jilin Prov. Inst. Ind. Health Occup. Dis.,
Jilin, Peop. Rep. China

SOURCE: Chem. Res. Toxicol. (1988), 1(5), 288-93
CODEN: CRTOEC

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Evaluation of the immunol. contribution to the pathogenesis of isocyanate lung disease necessitates prepn. of isocyanate-protein conjugates to detect anti-isocyanate antibodies. Sera were obtained from 2 guinea pigs immunized with MDI and from 3 workers with occupational exposure to MDI. By use of Western blot anal., guinea pig IgG antibodies were best detected by the monomeric component of MDI-guinea pig serum albumin. ELISA addnl. indicated that conjugates which contained a high d. of hapten detected greater amts. of antibody than did conjugates contg. low amts. of hapten. The same procedures were then used to assess the amt. of MDI-specific IgG and IgE antibody in sera from symptomatic workers. Effective MDI-HSA antigens were those that were monomeric and had high haptenic content. Highly substituted conjugates of monoisocyanates (Ph isocyanate and p-tolyl isocyanate) with serum albumins were also effective in detecting antibodies to MDI. These results indicate the importance of the compn. of isocyanate conjugate antigens in detecting antibodies to diisocyanates and suggest that stds. be developed for prepn. and characterization of these diagnostic serol. reagents.

L3 ANSWER 14 OF 14 CAPLUS COPYRIGHT 1999 ACS

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ACCESSION NUMBER: 1974:57087 CAPLUS
DOCUMENT NUMBER: 80:57087
TITLE: Modified procedure for evaluation of the
lecithin/sphingomyelin ratio in amniotic fluid
AUTHOR(S): Coch, Emily; Meyer, John S.; Goldman, Gordon;
Kessler, Gerald
CORPORATE SOURCE: Dep. Pathol. Lab. Med., Jew. Hosp., St. Louis,
Mo., USA
SOURCE: Clin. Chem. (1973), 19(9), 967-72
CODEN: CLCHAU
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Several modifications were made to the procedure of L. Gluck, et al. (1971) for evaluation of amniotic fluid lecithin/sphingomyelin (L/S) ratios. The acetone-pptn. step was eliminated, resulting in a faster extn. procedure and higher concns. of phospholipids. For faster thin-layer chromatog., silica gel-impregnated sheets of glass-fiber and a modified solvent system are used. For spot detection, a noncorrosive Bi subnitrate reagent specific for lecithin and sphingomyelin and an I vapor method to confirm the L/S ratio were used. The Bi spray also detected significant specimen contamination by phospholipids from plasma or erythrocytes. The L/S ratio was evaluated by visually comparing the relative size and color intensity of the lecithin and sphingomyelin TLC spots. An L/S ratio of 5 or more is consistent with mature pulmonary function and an L/S ratio of less than 2 suggests fetal pulmonary immaturity.

(FILE 'MEDLINE, BIOSIS, EMBASE, LIFESCI, WPIDS, CONFSCI, SCISEARCH, JICST-EPLUS, PROMT, CANCERLIT' ENTERED AT 12:12:56 ON 25 OCT 1999)

L4 210 S L3
L5 113 DUP REM L4 (97 DUPLICATES REMOVED)
L6 14 S L5 AND (HYBRIDIZ? OR HYBRIDIS?)

L6 ANSWER 1 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD

ACCESSION NUMBER: 1999-418749 [35] WPIDS
CROSS REFERENCE: 1998-414099 [35]; 1998-414100 [35]; 1998-414105
[35]; 1998-414114 [35]; 1998-427559 [35];
1998-506364 [42]; 1998-520811 [44]; 1998-609887
[42]; 1999-059865 [05]; 1999-080881 [07];
1999-120770 [10]; 1999-132229 [11]; 1999-132234
[11]; 1999-204988 [17]; 1999-430031 [36]

DOC. NO. CPI: C1999-123038

TITLE: New isolated human genes encoding secreted
polypeptides.

DERWENT CLASS: B04 D16

INVENTOR(S): CARTER, K C; DUAN, R D; FENG, P; FERRIE, A M;
FLORENCE, C; FLORENCE, K; GREENE, J M; JANAT, F;
KYAW, H; MOORE, P A; NI, J; ROSEN, C A; RUBEN, S M;
SHI, Y; SOPPET, D R; WEI, Y; YU, G

Searcher : Shears 308-4994



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PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 82
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG

WO 9931117	A1	19990624	(199935)*	EN	536
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM HR HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT					
LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL					
TJ TM TR TT UA UG US UZ VN YU ZW					

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE

WO 9931117	A1	WO 1998-US27059	19981217

PRIORITY APPLN. INFO: US 1997-68369 19971219; US 1997-68006
19971218; US 1997-68007 19971218; US
1997-68008 19971218; US 1997-68053
19971218; US 1997-68054 19971218; US
1997-68057 19971218; US 1997-68064
19971218; US 1997-70923 19971218; US
1997-68169 19971219; US 1997-68365
19971219; US 1997-68367 19971219; US
1997-68368 19971219

AN 1999-418749 [35] WPIDS
CR 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114
[35]; 1998-427559 [35]; 1998-506364 [42]; 1998-520811 [44];
1998-609887 [42]; 1999-059865 [05]; 1999-080881 [07]; 1999-120770
[10]; 1999-132229 [11]; 1999-132234 [11]; 1999-204988 [17];
1999-430031 [36]

AB WO 9931117 A UPAB: 19990908
NOVELTY - Isolated human nucleic acids (I) encoding secreted
proteins, are new.

DETAILED DESCRIPTION - (I) comprises a polynucleotide (PN)
having a nucleotide sequence (NS) at least 95% identical to:

(a) a PN fragment of one of a total of 110 defined human cDNA
sequences given in the specification or a PN fragment of the cDNA
sequence included in ATCC Deposit No. Z which is
hybridizable to one of the 110 defined cDNA sequences;

(b) a PN which is an (allelic) variant of one of the 110
defined cDNA sequences;

(c) a PN encoding a biologically active polypeptide or a
polypeptide fragment, domain or **epitope** of one of the 110

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defined amino acid sequences given in the specification or a polypeptide fragment, domain or **epitope**, respectively, encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridizable** to one of the defined cDNA sequences;

(d) a PN which encodes a species homolog of one of the 110 defined polypeptides; or

(e) a PN capable of **hybridizing** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridize** under stringent conditions to a sequence of only A residues or of only T residues.

ATCC Deposit No. Z refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are ATCC 209463, 209511, 209551.

INDEPENDENT CLAIMS are also included for the following:

- (1) a recombinant vector comprising (I);
- (2) a method of making a recombinant host cell comprising (I);
- (3) a recombinant host cell produced by a method as in (2);
- (4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 110 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z;
- (5) an isolated antibody that binds specifically to an isolated polypeptide as in (4);
- (6) a recombinant host cell that expresses an isolated polypeptide as above;
- (7) a method of making an isolated polypeptide by culturing the host cell of (6);
- (8) the polypeptide produced by the method of (7);
- (9) a gene corresponding to a cDNA sequence of the 110 defined amino acid sequences;
- (10) a method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
 - (a) determining the presence or absence of a mutation in (I); and
 - (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of the mutation;
- (11) a method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
 - (a) determining the presence or amount of expression of the polypeptide of (4) in a biological sample; and
 - (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide;
- (12) a method for identifying a binding partner to the polypeptide of (4) comprising:

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(a) contacting the polypeptide of (4) with a binding partner;
(b) determining whether the binding partner effects an activity of the polypeptide; and

(13) a method of identifying an activity in a biological assay, where the method comprises:

(a) expressing one of the 110 defined cDNAs in a cell;
(b) isolating the supernatant;
(c) detecting an activity in a biological assay; and
(d) identifying the protein in the supernatant having the activity; and

(14) the product produced by the method of (13).

ACTIVITY - Cytostatic; Nootropic; Neuroprotective; Osteopathic; Antiseborreic; Dermatological; Antipsoritic; Antidiabetic; Antiasthmatic.

MECHANISM OF ACTION - None given.

USE - The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be diagnosed by determining the amount of the new polypeptides in a sample or by determining the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 110 PNs, based on which tissues they are most highly expressed in, and include developing products for the diagnosis or treatment of cancer, tumors, neurodegenerative disorders, developmental abnormalities and fetal deficiencies, reproductive disorders, blood disorders, leukemias, diseases of the immune system, autoimmune diseases, hepatic and renal disease, lymphomas, inflammation, allergies, Alzheimer's and cognitive disorders, schizophrenia, disorders involving osteoclasts such as osteoporosis, arthritis, sepsis, acne, asthma, psoriasis, stroke, trauma, diseases of testes, lung or prostate, digestive/endocrine disorders, diabetes and AIDS. The polypeptides are also useful for identifying their binding partners (claimed).

Dwg.0/0

L6 ANSWER 2 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1999-303069 [25] WPIDS
CROSS REFERENCE: 1999-312869 [25]
DOC. NO. NON-CPI: N1999-227015
DOC. NO. CPI: C1999-089015
TITLE: New isolated human genes and the secreted polypeptides they encode.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): BREWER, L A; CARTER, K C; DUAN, D R; EBNER, R; ENDRESS, G A; FENG, P; FLORENCE, C; FLORENCE, K A; GREENE, J M; JANAT, F; KAYW, H; LAFLEUR, D W; MOORE, P A; NI, J; OLSEN, H S; ROSEN, C A; RUBEN, S M; SHI, Y; SOPPET, D R; WEI, Y; YOUNG, P
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC.

Searcher : Shears 308-4994

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COUNTRY COUNT: 83
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG

WO 9922243	A1	19990506	(199925)*	EN	546
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GD GE GH GM HR HU ID IL IS JP KE KG KP KR KZ LC LK LR LS					
LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK					
SL TJ TM TR TT UA UG US UZ VN YU ZW					
AU 9912734	A	19990517	(199939)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE

WO 9922243	A1	WO 1998-US22376	19981023
AU 9912734	A	AU 1999-12734	19981023

FILING DETAILS:

PATENT NO	KIND	PATENT NO

AU 9912734	A Based on	WO 9922243

PRIORITY APPLN. INFO: US 1997-63387 19971024; US 1997-62784
19971024; US 1997-63088 19971024; US
1997-63089 19971024; US 1997-63090
19971024; US 1997-63091 19971024; US
1997-63092 19971024; US 1997-63097
19971024; US 1997-63098 19971024; US
1997-63099 19971024; US 1997-63100
19971024; US 1997-63101 19971024; US
1997-63109 19971024; US 1997-63110
19971024; US 1997-63111 19971024; US
1997-63148 19971024; US 1997-63386 19971024

AN 1999-303069 [25] WPIDS

CR 1999-312869 [25]

AB WO 9922243 A UPAB: 19990707

NOVELTY - One hundred and forty eight isolated human genes and secreted proteins they encode are new.

DETAILED DESCRIPTION - An isolated nucleic acid molecule (NAM) (I) comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to:

(a) a PN fragment of one of a total of 148 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is

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hybridisable to one of the 148 defined cDNA sequence;

(b) a PN which is an (allelic) variant of one of the 148 defined cDNA sequences;

(c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or **epitope** of one of the 148 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequences;

(d) a PN which encodes a species homologue of one of the 148 defined polypeptides; or

(e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues.

INDEPENDENT CLAIMS are also included for:

(1) a recombinant vector comprising (I);

(2) a method of making a recombinant host cell comprising (I);

(3) a recombinant host cell produced by a method as in (2);

(4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 148 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z;

(5) an isolated antibody that binds specifically to an isolated polypeptide as in (4); and

(6) a recombinant host cell that expresses a gene corresponding to a cDNA sequence of the 148 defined amino acid sequences.

Note: From the disclosure 'ATCC Deposit No. Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are: ATCC 209299, 209346, 209300, 209324. an isolated polypeptide as above;

ACTIVITY - None given.

MECHANISM OF ACTION - None given.

USE - The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed** by **determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 148 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and fetal deficiencies, blood disorders, leukemias, diseases of the immune system, autoimmune diseases, hepatic and renal disease, lymphomas, inflammation, allergies, ischemic shock, Alzheimer's and cognitive

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disorders, schizophrenia, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, diseases of testes, lung or thymus, digestive/endocrine disorders, infections and AIDS. The polypeptides are also useful for identifying their binding partners (claimed).

L6 ANSWER 3 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1999-167452 [14] WPIDS
DOC. NO. CPI: C1999-048990
TITLE: New isolated human genes encoding secreted polypeptides - useful for diagnosis and treatment of pathological diseases.
DERWENT CLASS: B04 D16
INVENTOR(S): BREWER, L A; EBNER, R; FERRIE, A M; GREENE, J M; JANAT, F; NI, J; OLSEN, H S; ROSEN, C A; RUBEN, S M; SOPPET, D R; YOUNG, P E; YU, G
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 82
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9907891	A1	19990218	(199914)*	EN	329
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM HR HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT					
LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL					
TJ TM TR TT UA UG US UZ VN YU ZW					
AU 9887684	A	19990301	(199928)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9907891	A1	WO 1998-US16235	19980804
AU 9887684	A	AU 1998-87684	19980804

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9887684	A Based on	WO 9907891

PRIORITY APPLN. INFO: US 1997-56732 19970819; US 1997-54798
19970805; US 1997-54803 19970805; US
1997-54804 19970805; US 1997-54806
19970805; US 1997-54807 19970805; US
Searcher : Shears 308-4994

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1997-54808 19970805; US 1997-54809
19970805; US 1997-55309 19970805; US
1997-55310 19970805; US 1997-55311
19970805; US 1997-55312 19970805; US
1997-55386 19970805; US 1997-55970
19970818; US 1997-55986 19970818; US
1997-56364 19970819; US 1997-56365
19970819; US 1997-56366 19970819; US
1997-56367 19970819; US 1997-56370
19970819; US 1997-56371 19970819; US
1997-56557 19970819; US 1997-56563
19970819; US 1997-56731 19970819

AN 1999-167452 [14] WPIDS

AB WO 9907891 A UPAB: 19990412

NOVELTY - A total of 90 new isolated human genes encode secreted polypeptides which can be used in the diagnosis and treatment of pathological diseases such as cancers, neurological disorders, immune diseases, inflammation or blood disorders.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are included for the following: (1) An isolated nucleic acid molecule (NAM) (I) is claimed comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of one of a total of 90 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the 90 defined cDNA sequence; (b) a PN which is an (allelic) variant of one of the 90 defined cDNA sequences; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or **epitope** of one of the 90 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequences; (d) a PN which encodes a species homologue of one of the 90 defined polypeptides; or (e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues; (2) a recombinant vector comprising (I); (3) a method of making a recombinant host cell comprising (I); (4) a recombinant host cell produced by a method as in (3); (5) an isolated polypeptide comprising an amino acid sequence homologous to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 90 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z; (6) an isolated antibody that binds specifically to an isolated polypeptide as in (5); (7) a recombinant host cell that expresses an isolated polypeptide as above; and (8) a gene corresponding to a cDNA sequence of the 90 defined amino acid

Searcher : Shears 308-4994

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sequences. Note: From the disclosure 'ATCC Deposit No. Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are: ATCC 209146, 209177, 209179 and 209180.

USE - The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed** by **determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 90 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, CNS disorders, diseases of the immune system, autoimmune diseases, hepatic and renal disease, diabetes, inflammation, allergies, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, cardiovascular disorders, prostate diseases, asthma, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, **diseases** of testes, lung or thymus, digestive/endocrine disorders, infections and AIDS. The polypeptides are also useful for **identifying** their binding partners (claimed).

Dwg.0/0

L6 ANSWER 4 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1999-153691 [13] WPIDS
DOC. NO. CPI: C1999-045418
TITLE: New isolated human genes and the secreted polypeptides they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders.
DERWENT CLASS: B04 D16
INVENTOR(S): CARTER, K C; ENDRESS, G A; FAN, P; FENG, P; KYAW, H; LAFLEUR, D W; LI, Y; MOORE, P A; ROSEN, C A; RUBEN, S M; SHI, Y; WEI, Y; ZENG, Z
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 82
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG

WO 9906423	A1	19990211	(199913)*	EN	311
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM HR HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT					
LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL					
TJ TM TR TT UA UG US UZ VN YU ZW					

Searcher : Shears 308-4994

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AU 9887634 A 19990222 (199927)

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9906423	A1	WO 1998-US15949	19980729
AU 9887634	A	AU 1998-87634	19980729

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9887634	A Based on	WO 9906423

PRIORITY APPLN. INFO: US 1997-56730 19970819; US 1997-54209
19970730; US 1997-54211 19970730; US
1997-54212 19970730; US 1997-54213
19970730; US 1997-54214 19970730; US
1997-54215 19970730; US 1997-54217
19970730; US 1997-54218 19970730; US
1997-54234 19970730; US 1997-54236
19970730; US 1997-55968 19970818; US
1997-55969 19970818; US 1997-55972
19970818; US 1997-56534 19970819; US
1997-56543 19970819; US 1997-56554
19970819; US 1997-56561 19970819; US
1997-56727 19970819; US 1997-56729 19970819

AN 1999-153691 [13] WPIDS

AB WO 9906423 A UPAB: 19990331

An isolated nucleic acid molecule (NAM) (I) is claimed comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of one of a total of 83 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the 83 defined cDNA sequence; (b) a PN which is an (allelic) variant of one of the 83 defined cDNA sequences; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or **epitope** of one of the 83 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequences; (d) a PN which encodes a species homologue of one of the 83 defined polypeptides; or (e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues. Also claimed are: (1) a recombinant vector comprising (I); (2) a method of making a recombinant host cell

Searcher : Shears 308-4994



comprising (1); (3) a recombinant host cell produced by a method as in (2); (4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 83 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z; (5) an isolated antibody that binds specifically to an isolated polypeptide as in (4); (6) a recombinant host cell that expresses an isolated polypeptide as above; and (7) a gene corresponding to a cDNA sequence of the 83 defined amino acid sequences. Note: From the disclosure ATCC Deposit No. Z refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are: ATCC 209145, 209148, 209147.

USE- The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed** by **determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 83 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, leukemias, diseases of the immune system, autoimmune diseases, hepatic and renal disease, lymphomas, inflammation, allergies, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, restenosis, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, **diseases** of lung or skin, digestive/endocrine disorders, infections and AIDS. The polypeptides are also useful for **identifying** their binding partners (claimed).
Dwg.0/0

L6 ANSWER 5 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
 ACCESSION NUMBER: 1999-120770 [10] WPIDS
 CROSS REFERENCE: 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114 [35]; 1998-427559 [35]; 1998-506364 [42]; 1998-520811 [44]; 1998-609887 [42]; 1999-059865 [05]; 1999-080881 [07]; 1999-132229 [11]; 1999-132234 [11]; 1999-204988 [17]; 1999-418749 [32]; 1999-430031 [36]
 DOC. NO. CPI: C1999-035369
 TITLE: New isolated human genes and the secreted polypeptides they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders.
 Searcher : Shears 308-4994

09/092296

DERWENT CLASS: B04 D16
INVENTOR(S): BREWER, L A; EBNER, R; FISCHER, C L; KYAW, H;
LAPLEUR, D W; LI, Y; MOORE, P A; OLSEN, H S; ROSEN,
C A; RUBEN, S M; SHI, Y; SOPPET, D R; ZENG, Z
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 83
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9902546	A1	19990121	(199910)*	EN	462
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM GW HR HU ID IL IS JP KE KG KP KR KZ LC LK LR LS					
LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK					
SL TJ TM TR TT UA UG US UZ VN YU ZW					
AU 9884743	A	19990208	(199924)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9902546	A1	WO 1998-US13684	19980707
AU 9884743	A	AU 1998-84743	19980707

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9884743	A Based on	WO 9902546

PRIORITY APPLN. INFO: US 1997-58785 19970912; US 1997-51916
19970708; US 1997-51918 19970708; US
1997-51919 19970708; US 1997-51920
19970708; US 1997-51925 19970708; US
1997-51926 19970708; US 1997-51928
19970708; US 1997-51929 19970708; US
1997-51930 19970708; US 1997-51931
19970708; US 1997-51932 19970708; US
1997-52732 19970708; US 1997-52733
19970708; US 1997-52793 19970708; US
1997-52795 19970708; US 1997-52803
19970708; US 1997-55684 19970818; US
1997-55722 19970818; US 1997-55723
19970818; US 1997-55947 19970818; US
1997-55948 19970818; US 1997-55949
19970818; US 1997-55950 19970818; US
1997-55953 19970818; US 1997-55954
Searcher : Shears 308-4994

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19970818; US 1997-55964 19970818; US
1997-55984 19970818; US 1997-56360
19970818; US 1997-58660 19970912; US
1997-58661 19970912; US 1997-58664 19970912

AN 1999-120770 [10] WPIDS

CR 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114
[35]; 1998-427559 [35]; 1998-506364 [42]; 1998-520811 [44];
1998-609887 [42]; 1999-059865 [05]; 1999-080881 [07]; 1999-132229
[11]; 1999-132234 [11]; 1999-204988 [17]; 1999-418749 [32];
1999-430031 [36]

AB WO 9902546 A UPAB: 19990908

An isolated nucleic acid molecule (NAM) (I) is claimed comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of one of a total of 123 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the 123 defined cDNA sequence; (b) a PN which is an (allelic) variant of one of the 123 defined cDNA sequences; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or **epitope** of one of the 123 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequences; (d) a PN which encodes a species homologue of one of the 123 defined polypeptides; or (e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues. Also claimed are: (1) a recombinant vector comprising (I); (2) a method of making a recombinant host cell comprising (I); (3) a recombinant host cell produced by a method as in (2); (4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 123 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z; (5) an isolated antibody that binds specifically to an isolated polypeptide as in (4); (6) a recombinant host cell that expresses an isolated polypeptide as above; and (7) a gene corresponding to a cDNA sequence of the 123 defined amino acid sequences. Note: From the disclosure 'ATCC Deposit No. Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are: ATCC 209119, 209124, 209125, 209126.

USE - The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed by determining the**

Searcher : Shears 308-4994

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amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 123 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, leukemias, diseases of the immune system, autoimmune diseases, hepatic and renal disease, lymphomas, inflammation, allergies, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, restenosis, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, **diseases** of testes, lung or thyroid, digestive/endocrine disorders, infections and AIDS. The polypeptides are also useful for **identifying** their binding partners (claimed).

Dwg.0/0

L6 ANSWER 6 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1999-070066 [06] WPIDS
DOC. NO. CPI: C1999-020600
TITLE: New isolated human genes and the secreted polypeptides they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders.
DERWENT CLASS: B04 D16
INVENTOR(S): BREWER, L A; DUAN, R; EBNER, R; FERRIE, A M; FLORENCE, K A; GREENE, J M; HU, J; LAFLEUR, D W; MOORE, P A; NI, J; OLSEN, H S; ROSEN, C A; RUBEN, S M; SHI, Y; YOUNG, P
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 81
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9842738	A1	19981001	(199906)*	EN	385
RW: AT BE CH DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZW					
AU 9865646	A	19981020	(199909)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9842738	A1	WO 1998-US5311	19980319
		Searcher :	Shears 308-4994

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AU 9865646 A

AU 1998-65646 19980319

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9865646	A Based on	WO 9842738

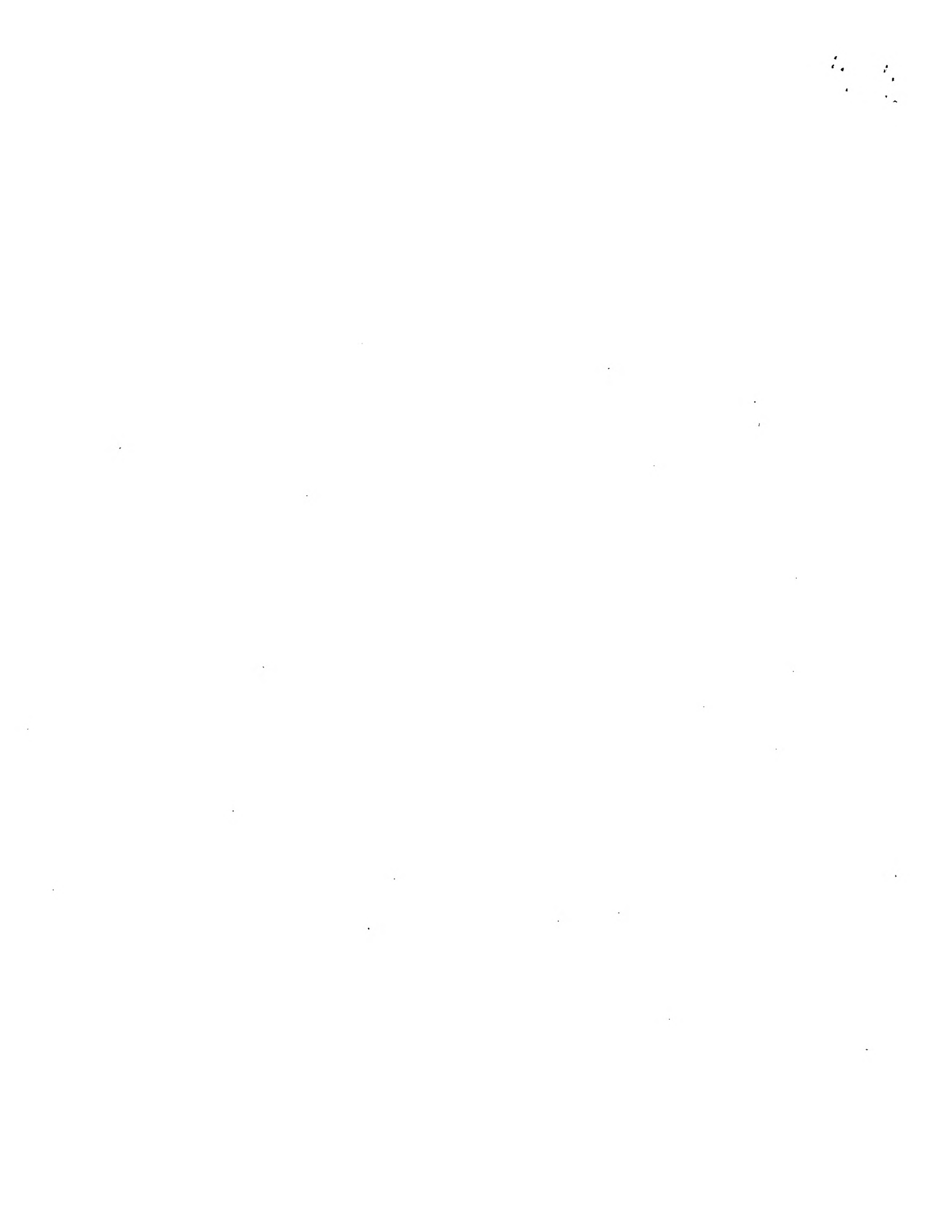
PRIORITY APPLN. INFO: US 1997-50937 19970530; US 1997-41276
19970321; US 1997-41277 19970321; US
1997-41281 19970321; US 1997-42344
19970321; US 1997-48069 19970530; US
1997-48094 19970530; US 1997-48095
19970530; US 1997-48096 19970530; US
1997-48099 19970530; US 1997-48131
19970530; US 1997-48135 19970530; US
1997-48154 19970530; US 1997-48160
19970530; US 1997-48186 19970530; US
1997-48187 19970530; US 1997-48188
19970530; US 1997-48351 19970530; US
1997-48352 19970530; US 1997-48355 19970530

AN 1999-070066 [06] WPIDS

AB WO 9842738 A UPAB: 19990210

An isolated nucleic acid molecule (NAM) (I) comprises a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of one of a total of 87 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the 87 defined cDNA sequences; (b) a PN which is an (allelic) variant of one of the 87 defined cDNA sequences; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or **epitope** of one of the 87 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequence; (d) a PN which encodes a species homologue of one of the 87 defined polypeptides; or (e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues. Also claimed are: (1) a recombinant vector comprising (I); (2) a method of making a recombinant host cell comprising (I); (3) a recombinant host cell produced by a method as in (2); (4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 87 defined amino acid sequences or the encoded sequence included in ATCC

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Deposit No. Z; (5) an isolated antibody that binds specifically to an isolated polypeptide as in (4); (6) a recombinant host cell that expresses an isolated polypeptide as in (4); and (7) a gene corresponding to a cDNA sequence of the 87 defined amino acid sequences. Note: From the disclosure 'ATCC Deposit No. Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are: ATCC 97923, 97924, 97957, 97958, 209071, 209641, 209072 and 209073.

USE - The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed by determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 87 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, leukaemias, diseases of the immune system (including allergies or asthma), lymphocytic diseases, brain associated diseases, hepatic and renal disease, lymphomas, inflammation, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, restenosis, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, **diseases** of testis, lung or thymus, digestive/endocrine disorders, including metabolic regulation, malabsorption, gastritis and neoplasms, and AIDS. The polypeptides are also useful for **identifying** their binding partners (claimed).

Dwg.0/0

L6 ANSWER 7 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1999-060335 [05] WPIDS
DOC. NO. NON-CPI: N1999-044750
DOC. NO. CPI: C1999-018022
TITLE: New LS170 nucleic acid from lung tissue - useful
for **detecting**, monitoring, preventing and
treating lung **disease**,
particularly cancer.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): BILLING-MEDEL, P A; COHEN, M; COLPITTS, T L;
FRIEDMAN, P N; GORDON, J; GRANADOS, E N; HODGES, S
C; KLASS, M R; KRATOCHVIL, J D; ROBERTS-RAPP, L;
RUSSELL, J C; STROUPE, S D
PATENT ASSIGNEE(S): (ABBO) ABBOTT LAB
COUNTRY COUNT: 20
PATENT INFORMATION:

Searcher : Shears 308-4994

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PATENT NO	KIND	DATE	WEEK	LA	PG

WO 9856951	A1	19981217	(199905)*	EN	119
RW: AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE					
W: CA JP					

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE

WO 9856951	A1	WO 1998-US11601	19980611

PRIORITY APPLN. INFO: US 1997-49183 19970611

AN 1999-060335 [05] WPIDS

AB WO 9856951 A UPAB: 19990210

Detection of a target LS170 polynucleotide (I) comprises treating a test sample with at least one LS170-specific nucleic acid (II) that has at least 50% identity with any of 9 sequences ((1)-(9), reproduced, of 251, 243, 258, 202, 458, 273, 200,1009 and 1027 bp, respectively), their fragments or complements. Also new are (a) LS170 sequences, or their fragments, that **hybridise** selectively to the LS170 gene and have at least 50% identity with (i) (1)-(9), other than (5) or their complements, or (ii) fragments of (1)-(7); (b) recombinant expression system containing (II) plus a control sequence; (c) cells transformed with this expression system; (d) LS170 polypeptides (III) at least 50% identical with 9 sequences ((23)-(31), of 256, 18, 26, 29, 12, 15, 11 or 19 amino acids (aa), reproduced), or their fragments, containing at least one LS170 **epitope**; (e) antibodies (Ab) specific for at least one LS170 **epitope** present on (III); (f) cells transformed with any of (1)-(9), their fragments or complements, encoding at least one **epitope** of LS170; (g) (I) having at least 50% identity with (i) any of (1)-(9), their fragments or complements or (ii) fragments of (1)-(7); (h) genes, or their fragments, encoding LS170 proteins at least 50% identical with (23); (j) genes, or their fragments, at least 50% identical with (8) or (9).

USE - LS170 represents a set of contiguous, partially overlapping sequences transcribed from lung tissue. They are used for **diagnosis**, staging, monitoring, in vivo imaging, prevention and treatment of **lung disease**, specifically cancer, and to indicate predisposition to such disease. Particularly **detection** of (I), LS170 antigens (using Ab in immunoassays) or anti-LS170 antibodies (using LS170 as antigen) is indicative of disease. Cells of (c) are used to express recombinant (III), used to raise Ab and for drug screening. LS170-related nucleic acid can be used to isolate related sequences; as standards and **reagents** in assays; as targets for drug screening, and as components or targets for therapy, e.g. as antisense, ribozyme or

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triplex-forming agents. Ab can be used to deliver therapeutic agents to LS170-expressing cells; directly as therapeutic agents (by neutralising LS170 polypeptides); in competitive binding drug screens, and to generate anti-idiotypic antibodies for use in rational drug design.

Dwg.0/4

L6 ANSWER 8 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1999-059865 [05] WPIDS
CROSS REFERENCE: 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114 [35]; 1998-427559 [35]; 1998-506364 [42]; 1998-520811 [44]; 1998-609887 [42]; 1999-080881 [07]; 1999-120770 [10]; 1999-132229 [11]; 1999-132234 [11]; 1999-204988 [17]; 1999-418749 [32]; 1999-430031 [36]
DOC. NO. NON-CPI: N1999-044486
DOC. NO. CPI: C1999-017701
TITLE: New isolated human genes and the secreted polypeptides they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): BREWER, L A; CARTER, K C; DILLON, P J; EBNER, R; ENDRESS, G A; FAN, P; FENG, P; FERRIE, A M; FISCHER, C L; FLORENCE, C; FLORENCE, K; GREENE, J M; HU, J; KYAW, H; LAFLEUR, D W; LI, Y; MOORE, P A; NI, J; OLSEN, H S; ROSEN, C A; RUBEN, S M; SHI, Y; SOPPET, D R; WEI, Y; YOUNG, P; YU, G; ZENG, Z
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 82
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG

WO 9854963	A2	19981210	(199905)*	EN	770
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT					
LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL					
TJ TM TR TT UA UG US UZ VN YU ZW					
AU 9878120	A	19981221	(199919)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE

WO 9854963	A2	WO 1998-US11422	19980604
AU 9878120	A	AU 1998-78120	19980604
Searcher		:	Shears 308-4994

09/092296

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9878120	A Based on	WO 9854963

PRIORITY APPLN. INFO: US 1997-70923 19971218; US 1997-48875
19970606; US 1997-48876 19970606; US
1997-48877 19970606; US 1997-48878
19970606; US 1997-48880 19970606; US
1997-48881 19970606; US 1997-48882
19970606; US 1997-48883 19970606; US
1997-48884 19970606; US 1997-48885
19970606; US 1997-48892 19970606; US
1997-48893 19970606; US 1997-48894
19970606; US 1997-48895 19970606; US
1997-48896 19970606; US 1997-48897
19970606; US 1997-48898 19970606; US
1997-48899 19970606; US 1997-48900
19970606; US 1997-48901 19970606; US
1997-48915 19970606; US 1997-48916
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1997-48949 19970606; US 1997-48962
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1997-48964 19970606; US 1997-48970
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1997-48972 19970606; US 1997-48974
19970606; US 1997-49019 19970606; US
1997-49020 19970606; US 1997-49373
19970606; US 1997-49374 19970606; US
1997-49375 19970606; US 1997-57584
19970905; US 1997-57627 19970905; US
1997-57628 19970905; US 1997-57629
19970905; US 1997-57634 19970905; US
1997-57635 19970905; US 1997-57642
19970905; US 1997-57643 19970905; US
1997-57644 19970905; US 1997-57645
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1997-57647 19970905; US 1997-57648
19970905; US 1997-57649 19970905; US
1997-57650 19970905; US 1997-57651
19970905; US 1997-57654 19970905; US
1997-57661 19970905; US 1997-57662
19970905; US 1997-57666 19970905; US
1997-57667 19970905; US 1997-57668
19970905; US 1997-57760 19970905; US
1997-57761 19970905; US 1997-57762
19970905; US 1997-57763 19970905; US
Searcher : Shears 308-4994

1997-57764 19970905; US 1997-57765
 19970905; US 1997-57769 19970905; US
 1997-57770 19970905; US 1997-57771
 19970905; US 1997-57774 19970905; US
 1997-57775 19970905; US 1997-57776
 19970905; US 1997-57777 19970905; US
 1997-57778 19970905

AN 1999-059865 [05] WPIDS

CR 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114
 [35]; 1998-427559 [35]; 1998-506364 [42]; 1998-520811 [44];
 1998-609887 [42]; 1999-080881 [07]; 1999-120770 [10]; 1999-132229
 [11]; 1999-132234 [11]; 1999-204988 [17]; 1999-418749 [32];
 1999-430031 [36]

AB WO 9854963 A UPAB: 19990908

An isolated nucleic acid molecule (NAM) (I) comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of one of a total of 207 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the 207 defined cDNA sequence; (b) a PN which is an (allelic) variant of one of the 207 defined cDNA sequences; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or epitope of one of the 207 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequences; (d) a PN which encodes a species homologue of one of the 207 defined polypeptides; or (e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues. Also claimed are: (1) a recombinant vector comprising (I); (2) a method of making a recombinant host cell comprising (I); (3) a recombinant host cell produced by a method as in (2); (4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 207 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z; (5) an isolated antibody that binds specifically to an isolated polypeptide as in (4); (6) a recombinant host cell that expresses an isolated polypeptide as above; and (7) a gene corresponding to a cDNA sequence of the 207 defined amino acid sequences.

Note: From the disclosure 'ATCC Deposit No. Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are: ATCC 97979, 97974, 97975, 97976, 97977, 209007, 209008,

Searcher : Shears 308-4994

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209009, 209010, 209011, 209080, 209081, 209082, 209083, 209084,
209085, 209511,.

USE The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed by determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 207 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, leukemias, diseases of the immune system, autoimmune diseases, hepatic and renal disease, lymphomas, inflammation, allergies, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, restenosis, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, **diseases of testes, lung or** thymus, digestive/endocrine disorders, infections and AIDS. The polypeptides are also useful for **identifying** their binding partners (claimed).

Dwg.0/0

L6 ANSWER 9 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1998-609887 WPIDS
CROSS REFERENCE: 1998-414099 [35]; 1998-414100 [35]; 1998-414105
[35]; 1998-414114 [35]; 1998-427559 [35];
1998-506364 [42]; 1998-520811 [44]; 1999-059865
[05]; 1999-080881 [07]; 1999-120770 [10];
1999-132229 [11]; 1999-132234 [11]; 1999-204988
[17]; 1999-418749 [32]; 1999-430031 [36]
DOC. NO. NON-CPI: N1998-474475
DOC. NO. CPI: C1998-182720
TITLE: New isolated human genes and the secreted
polypeptides they encode - useful for diagnosis and
treatment of e.g. cancers, neurological disorders,
immune diseases, inflammation or blood disorders.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): BEDNARIK, D P; BREWER, L A; CARTER, K C; DUAN, R;
EBNER, R; ENDRESS, G A; FENG, P; FERRIE, A M;
FISCHER, C L; FLORENCE, K A; GREENE, J M; HU, J;
NI, J; OLSEN, H S; ROSEN, C A; RUBEN, S M; SOPPET,
D R; YOUNG, P E; YU, G; KYAW, H; LAFLEUR, D W; LI,
Y; MOORE, P A; SHI, Y; ZENG, Z
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 81
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
	Searcher	:	Shears	308-4994	

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WO 9839446 A2 19980911 (199851)* EN 447
RW: AT BE CH DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW
NL OA PT SD SE SZ UG ZW
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI
GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT
LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL
TJ TM TR TT UA UG US UZ VN YU ZW
AU 9865452 A 19980922 (199908)

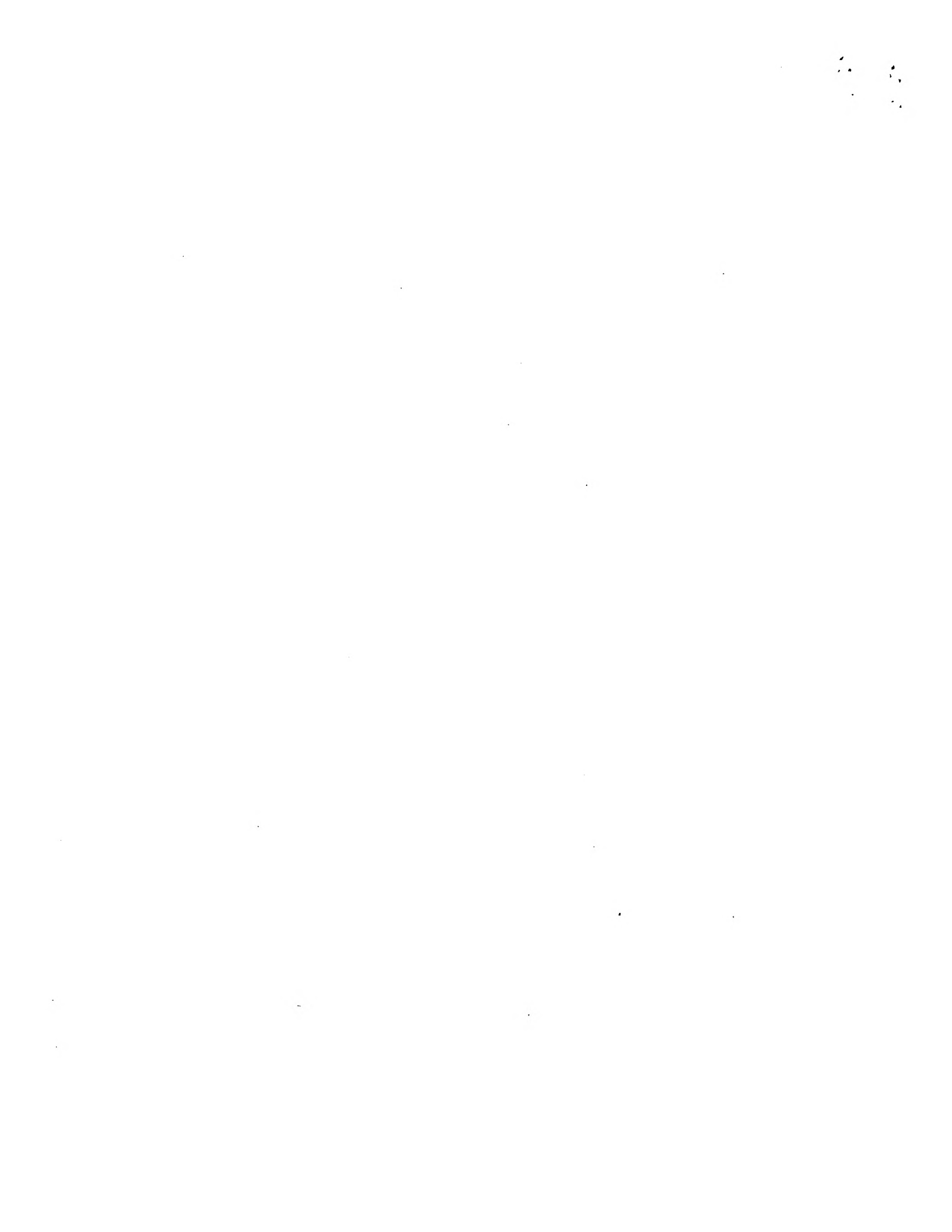
APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9839446	A2	WO 1998-US4482	19980306
AU 9865452	A	AU 1998-65452	19980306

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9865452	A Based on	WO 9839446

PRIORITY APPLN. INFO: US 1997-57761 19970905; US 1997-38621
19970307; US 1997-40161 19970307; US
1997-40162 19970307; US 1997-40163
19970307; US 1997-40333 19970307; US
1997-40334 19970307; US 1997-40336
19970307; US 1997-40626 19970307; US
1997-43311 19970411; US 1997-43312
19970411; US 1997-43313 19970411; US
1997-43314 19970411; US 1997-43315
19970411; US 1997-43568 19970411; US
1997-43569 19970411; US 1997-43576
19970411; US 1997-43578 19970411; US
1997-43580 19970411; US 1997-43669
19970411; US 1997-43670 19970411; US
1997-43671 19970411; US 1997-43672
19970411; US 1997-43674 19970411; US
1997-47492 19970523; US 1997-47500
19970523; US 1997-47501 19970523; US
1997-47502 19970523; US 1997-47503
19970523; US 1997-47581 19970523; US
1997-47582 19970523; US 1997-47583
19970523; US 1997-47584 19970523; US
1997-47585 19970523; US 1997-47586
19970523; US 1997-47587 19970523; US
1997-47588 19970523; US 1997-47589
19970523; US 1997-47590 19970523; US
Searcher : Shears 308-4994



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1997-47592 19970523; US 1997-47593
19970523; US 1997-47594 19970523; US
1997-47595 19970523; US 1997-47596
19970523; US 1997-47597 19970523; US
1997-47598 19970523; US 1997-47599
19970523; US 1997-47600 19970523; US
1997-47601 19970523; US 1997-47612
19970523; US 1997-47613 19970523; US
1997-47614 19970523; US 1997-47615
19970523; US 1997-47617 19970523; US
1997-47618 19970523; US 1997-47632
19970523; US 1997-47633 19970523; US
1997-48964 19970606; US 1997-48974
19970606; US 1997-56630 19970822; US
1997-56631 19970822; US 1997-56632
19970822; US 1997-56636 19970822; US
1997-56637 19970822; US 1997-56662
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1997-56845 19970822; US 1997-56862
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1997-56872 19970822; US 1997-56874
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1997-56876 19970822; US 1997-56877
19970822; US 1997-56878 19970822; US
1997-56879 19970822; US 1997-56880
19970822; US 1997-56881 19970822; US
1997-56882 19970822; US 1997-56884
19970822; US 1997-56886 19970822; US
1997-56887 19970822; US 1997-56888
19970822; US 1997-56889 19970822; US
1997-56892 19970822; US 1997-56893
19970822; US 1997-56894 19970822; US
1997-56903 19970822; US 1997-56908
19970822; US 1997-56909 19970822; US
1997-56910 19970822; US 1997-56911
19970822; US 1997-57650 19970905

AN 1998-609887 WPIDS

CR 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114
[35]; 1998-427559 [35]; 1998-506364 [42]; 1998-520811 [44];
1999-059865 [05]; 1999-080881 [07]; 1999-120770 [10]; 1999-132229
[11]; 1999-132234 [11]; 1999-204988 [17]; 1999-418749 [32];
1999-430031 [36]

AB WO 9839446 A UPAB: 19990908

An isolated nucleic acid molecule (NAM) (I) comprising a
polynucleotide (PN) having a nucleotide sequence (NS) at least 95%
identical to:

(a) a PN fragment of one of a total of 70 defined human cDNA
sequences given in the specification or a PN fragment of the cDNA
sequence included in ATCC Deposit No. Z, which is

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hybridisable to one of the 70 defined cDNA sequence;

(b) a PN which is an (allelic) variant of one of the 70 defined cDNA sequences;

(c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or **epitope** of one of the 70 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequence;

(d) a PN which encodes a species homologue of one of the 70 defined polypeptides., or

(e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues.

Also claimed are:

(1) a recombinant vector comprising (I),

(2) a method of making a recombinant host cell comprising (1);

(3) a recombinant host cell produced by a method as in (2);

(4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 70 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z;

(5) an isolated antibody that binds specifically to an isolated polypeptide as in (4);

(6) a recombinant host cell that expresses an isolated polypeptide as in (4), and

(7) a gene corresponding to a cDNA sequence of the 70 defined amino acid sequences.

Note., From the disclosure 'ATCC Deposit No. Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are- ATCC 97900, 97901, 97903, 97898, 209010, 97897, 97899, 97897, 97976, 97904, 97926, 209043-209050, 209085, 209084, 209236.

USE - The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed** by **determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 70 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, leukaemias, diseases of the immune system (including allergies or asthma), lymphocytic diseases, brain associated diseases, hepatic and renal disease, lymphomas,

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inflammation, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, restenosis, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, diseases of testis, lung or thymus, digestive/endocrine disorders, including metabolic regulation, malabsorption, gastritis and neoplasms, and AIDS. The polypeptides are also useful for identifying their binding partners (claimed).

Dwg.0/10

L6 ANSWER 10 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1998-594496 [50] WPIDS
DOC. NO. NON-CPI: N1998-462620
DOC. NO. CPI: C1998-178291
TITLE: New isolated human genes and secreted polypeptide(s) they encode - useful for the diagnosis and treatment of e.g. cancers, CNS disorders, immune system disorders, inflammatory disease and bacterial infections.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): FENG, P; NI, J; ROSEN, C A; RUBEN, S M; YU, G
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 82
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG

WO 9845712	A2	19981015	(199850)*	EN	141
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT					
LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL					
TJ TM TR TT UA UG US UZ VN YU ZW					
AU 9869529	A	19981030	(199911)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE

WO 9845712	A2	WO 1998-US6801	19980407
AU 9869529	A	AU 1998-69529	19980407

FILING DETAILS:

PATENT NO	KIND	PATENT NO

AU 9869529	A Based on	WO 9845712

Searcher : Shears 308-4994

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PRIORITY APPLN. INFO: US 1997-48184 19970530; US 1997-42726
 19970408; US 1997-42727 19970408; US
 1997-42728 19970408; US 1997-42754
 19970408; US 1997-42825 19970408; US
 1997-48068 19970530; US 1997-48070 19970530

AN 1998-594496 [50] WPIDS

AB WO 9845712 A UPAB: 19990210

An isolated nucleic acid molecule (NAM) (I) is claimed comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of a total of 20 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the 20 defined cDNA sequences; (b) a PN which is an (allelic) variant of one of the 20 defined cDNA sequence; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or **epitope** of one of the 20 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is **hybridisable** to one of the defined cDNA sequences; (d) a PN which encodes a species homologue of one of the 20 defined polypeptides; or (e) a PN capable of **hybridising** under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not **hybridise** under stringent conditions to a sequence of only A residues or of only T residues. Also claimed are: (1) a recombinant vector comprising (I); (2) a method of making a recombinant host cell comprising (I); (3) a recombinant host cell produced by a method as in (2); (4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, **epitope**, secreted form, full-length protein, (allelic) variant or species homologue of one of the 20 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z; (5) an isolated antibody that binds specifically to an isolated polypeptide as in (4); (6) a recombinant host cell that expresses an isolated polypeptide as in (4); (7) a gene corresponding to a cDNA sequence of one of the 20 nucleotide sequences given in the specification. Note: From the disclosure "ATCC deposit No. Z" refers to ATCC No's 97955 and 209074.

USE - The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be **diagnosed** by **determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 20 PNs, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of central nervous system (CNS) and immune system diseases, reproductive disorders, cancers,

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congenital malformations, degenerative diseases, trauma, inflammatory disease, neoplasia, metabolic disorders, diseases in testes, placenta, liver, brain and activated T cells, spleen diseases, lung diseases, heart diseases, rhabdomyosarcoma and disorders of the endocrine system or other endocrinopathies, e.g. endocrine polyglandular syndrome, endocrinoma, and endocrine ophthalmopathy, osteoclastoma and other bone remodelling disorders, bacterial infections and sepsis. The polypeptides are also useful for identifying their binding partners (claimed).
Dwg.0/0

L6 ANSWER 11 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1998-506364 [43] WPIDS
CROSS REFERENCE: 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114 [35]; 1998-427559 [35]; 1998-520811 [44]; 1998-609887 [42]; 1999-059865 [05]; 1999-080881 [07]; 1999-120770 [10]; 1999-132229 [11]; 1999-132234 [11]; 1999-204988 [17]; 1999-418749 [32]; 1999-430031 [36]
DOC. NO. NON-CPI: N1998-394741
DOC. NO. CPI: C1998-152795
TITLE: New isolated human genes and the secreted polypeptide(s) they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): BEDNARIK, D P; BREWER, L A; CARTER, K C; DUAN, R; EBNER, R; ENDRESS, G A; FENG, P; FERRIE, A M; FISCHER, C L; FLORENCE, K A; GREENE, J M; HU, J; NI, J; OLSEN, H S; ROSEN, C A; RUBEN, S M; SOPPET, D R; YOUNG, P E; YU, G; KYAW, H; LAFLEUR, D W; LI, Y; MOORE, P A; SHI, Y; ZENG, Z
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 81
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG

WO 9839448	A2	19980911	(199843)*	EN	721
RW: AT BE CH DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW					
NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT					
LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL					
TJ TM TR TT UA UG US UZ VN YU ZW					
AU 9865453	A	19980922	(199908)		
AU 9891304	A	19990322	(199931)		

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APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9839448	A2	WO 1998-US4493	19980306
AU 9865453	A	AU 1998-65453	19980306
AU 9891304	A	AU 1998-91304	19980903

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9865453	A Based on	WO 9839448
AU 9891304	A Based on	WO 9911293

PRIORITY APPLN. INFO: US 1997-61060 19971002; US 1997-38621
19970307; US 1997-40161 19970307; US
1997-40162 19970307; US 1997-40163
19970307; US 1997-40333 19970307

AN 1998-506364 [43] WPIDS
CR 1998-414099 [35]; 1998-414100 [35]; 1998-414105 [35]; 1998-414114
[35]; 1998-427559 [35]; 1998-520811 [44]; 1998-609887 [42];
1999-059865 [05]; 1999-080881 [07]; 1999-120770 [10]; 1999-132229
[11]; 1999-132234 [11]; 1999-204988 [17]; 1999-418749 [32];
1999-430031 [36]

AB WO 9839448 A UPAB: 19990908
An isolated nucleic acid molecule (I) comprises a polynucleotide
(PN) having a nucleotide sequence at least 95% identical to: (a) a
PN fragment of one of a total of 186 defined human cDNA sequences
given in the specification or a PN fragment of the cDNA sequence
included in ATCC Deposit No: Z, which is **hybridisable** to
one of the 186 defined cDNA sequences; (b) a PN which is an
(allelic) variant of one of the 186 defined cDNA sequences; (c) a PN
encoding a biologically active polypeptide or a polypeptide
fragment, domain or **epitope** of one of the 186 defined
amino acid sequences given in the specification or a polypeptide
fragment encoded by a cDNA sequence included in ATCC Deposit No. Z,
which is **hybridisable** to one of the defined cDNA
sequences; (d) a PN which encodes a species homologue of one of the
186 defined polypeptides; or (e) a PN capable of **hybridising**
under stringent conditions to any one of the PNs specified in
(a)-(d), where the PN does not **hybridise** under stringent
conditions to a sequence of only A residues or of only T residues.

Also claimed is an isolated polypeptide comprising an amino
acid sequence at least 95% identical to a sequence selected from:
(a) a polypeptide fragment (preferably having biological activity),
domain, **epitope**, secreted form, full-length protein,
(allelic) variant or species homologue of one of the 186 defined

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amino acid sequences or the encoded sequence included in ATCC
Deposit No: Z.

Note: From the disclosure 'ATCC Deposit No: Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the American Type Culture Collection. The deposit numbers are: ATCC 97897-97904 and ATCC 209076, 209215, 209139, 209011, 209119, 209235, 209627 and 209043-209050.

USE - The polynucleotides and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also, pathological conditions can be **diagnosed** by **determining** the amount of the new polypeptides in a sample or by **determining** the presence of mutations in the new polynucleotides (claimed). Specific uses are described for each of the 186 polynucleotides, based on which tissues they are most highly expressed in, and include developing products for the **diagnosis** or treatment of: cancer; tumours; neurodegenerative disorders; developmental abnormalities and foetal deficiencies; blood disorders; leukaemias; diseases of the immune system (including allergies or asthma); lymphocytic diseases; brain associated diseases; hepatic and renal disease; lymphomas; inflammation; ischaemic shock; Alzheimer's and cognitive disorders; schizophrenia; restenosis; prostate diseases; obesity; disorders involving osteoclasts such as osteoporosis, arthritis or malignancies; **diseases** of testis, **lung** or thymus; thyroiditis and thyroid tumours; digestive/endocrine disorders, including metabolic regulation, malabsorption, gastritis and neoplasms; and AIDS. The polypeptides are also useful for **identifying** their binding partners (claimed).

Dwg.0/0

L6 ANSWER 12 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1998-437479 [37] WPIDS
DOC. NO. NON-CPI: N1998-340776
DOC. NO. CPI: C1998-133112
TITLE: New nucleic acid for the **lung**
disease marker LU105 - polypeptides,
antibodies and genes, used for **diagnosis**,
prevention, treatment of **lung**
disease, specifically cancer.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): BILLING-MEDEL, P A; COHEN, M; COLPITTS, T L;
FRIEDMAN, P N; GORDON, J; GRANADOS, E N; HODGES, S
C; KLASS, M R; KRATOCHVIL, J D; ROBERTSRAPP, L;
RUSSELL, J C; STROUPE, S D
PATENT ASSIGNEE(S): (ABBO) ABBOTT LAB
COUNTRY COUNT: 19
PATENT INFORMATION:

Searcher : Shears 308-4994

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PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9833926	A1	19980806	(199837)*	EN	117
RW: AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE					
W: CA JP					

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9833926	A1	WO 1998-US1766	19980130

PRIORITY APPLN. INFO: US 1997-791710 19970131

AN 1998-437479 [37] WPIDS

AB WO 9833926 A UPAB: 19980916

A method for detecting target LU105 nucleic acid (I) comprises treating a sample with at least one LU105-specific nucleic acid (II), or its complement. (II) is at least 50% identical with 190, 244, 225, 114, 562 or 519 bp sequences given in the specification, or their fragments and complements.

Also claimed are:

(1) (I) or its fragments able to **hybridise** selectively to the LU105 gene and having at least 50% identity with the 190, 244, 225, 114, 562 or 519 bp sequences given above;

(2) recombinant expression systems including (I) and control sequence;

(3) cells transformed with this expression system;

(4) LU105 polypeptides (III) at least 50% identical with the 104, 26, 19, 21, 18 or 19 amino acid (aa) sequences given in the specification or their fragments;

(5) antibodies (Ab) that bind to at least one LU105 **epitope** present in (III);

(6) cells transformed with the 190, 244, 225, 114, 562 or 519 bp sequences described above;

(7) LU105 specific nucleic acid (II); and

(8) genes, or their fragments, that encode a protein at least 50% identical with the 104 aa sequences as in (4).

USE - LU105 is a **lung disease** marker. Cells as in (3) are used to express recombinant (III) which are used to raise Ab. Ab are used to **detect** the LU105 antigen, and correspondingly this antigen is used to **detect** specific antibodies, in usual immunoassays. (I) and (III) are used for **diagnosis**, staging, monitoring, prognosis, prevention, treatment (e.g. using antisense molecules, ribozymes, Ab or other antagonists) and **determination** of susceptibility to, **lung disease**, specifically cancer. (III) are also used to screen for specific binding agents, potentially useful

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therapeutically. LU105 is a marker for lung disease (present at high concentration, in altered form or in an unusual body compartment).

ADVANTAGE - LU105 can be detected in blood, plasma or serum in an inexpensive, non-invasive test.

Dwg.0/6

L6 ANSWER 13 OF 14 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD
ACCESSION NUMBER: 1998-333327 [29] WPIDS
DOC. NO. NON-CPI: N1998-260122
DOC. NO. CPI: C1998-103375
TITLE: Human chemokine beta-13 polypeptide - useful in diagnosis and treatment of immune-system related disorders e.g. cancer of the immune system, leukaemias, autoimmune diseases etc..
DERWENT CLASS: B04 D16 S03
INVENTOR(S): LI, H; SEIBEL, G
PATENT ASSIGNEE(S): (HUMA-N) HUMAN GENOME SCI INC
COUNTRY COUNT: 79
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9824908	A1	19980611	(199829)*	EN	86
RW: AT BE CH DE DK EA ES FI FR GB GH GR IE IT KE LS LU MC MW NL					
OA PT SD SE SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV					
MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM					
TR TT UA UG US UZ VN YU ZW					
AU 9853834	A	19980629	(199845)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9824908	A1	WO 1997-US23144	19971205
AU 9853834	A	AU 1998-53834	19971205

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9853834	A Based on	WO 9824908

PRIORITY APPLN. INFO: US 1996-32432 19961205
AN 1998-333327 [29] WPIDS
AB WO 9824908 A UPAB: 19980722
Isolated human chemokine beta -13 (CK beta -13) polypeptide
Searcher : Shears 308-4994



comprises amino acid sequence: (a) 95 % identical or more to 93 amino acid sequence (I) given in the specification, (b) mature CK beta -13 polypeptide sequence comprising amino acids 25-93 or 29-93 of sequence (I); (c) encoded by cDNA clone contained in ATCC 97113; or (d) a sequence complementary to the sequences as in (a)-(b).

Also claimed are: (1) polypeptide comprising **epitope**-bearing portion of CK beta -13 protein comprising amino acids Thr22-Gly28, Asn30-Leu47, Thr56-Val65 or Phe70-Trp83 of sequence (I); (2) nucleic acid molecules as follows: (i) encoding CK beta -13 polypeptide sequences as above; (ii) complementary to (i); (iii) **hybridising** to (i) but not to polynucleotide with sequence consisting of only A or only T residues; (iv) encoding **epitope**-bearing portion of (i), optionally as in (1); (v) having 282 bp sequence (II) and optionally encoding CK beta -13 polypeptide sequences as above; (vi) at least 95 % identical to sequence encoding polypeptide with amino acids n-93, 1-m and n-m (n and m are integers between 1-35 and 77-93 respectively) of (i); and (vii) encoding polypeptide which is a portion of CK beta -13 sequence encoded by cDNA clone in ATCC 97113 excluding 1-35 amino acids from amino terminus and/or 1-17 amino acids from carboxy terminus; (3) recombinant vectors produced by inserting (i) into vector; (4) host cells produced by introducing the vector as in (4); and (5) antibody binding specifically to CK beta -13 polypeptide.

USE - The polypeptides and nucleic acids are useful in **diagnosis** and treatment of immune system-related disorders in mammals (preferably humans). Such disorders include tumours, cancers, **interstitial lung disease** and disregulation of immune cell function including leukaemias, lymphomas, autoimmune diseases etc. For example, certain tissues in mammals with cancer of the immune system express enhanced/decreased levels of CK beta -13 and mRNA encoding CK beta -13, and **diagnosis** can be achieved by assaying CK beta -13 gene expression and comparing to standard levels. The polypeptides can be administered therapeutically in pharmaceutical compositions e.g. to treat immune system-related disorders as above, treat sepsis, inhibit bone marrow stem cell colony formation during cancer therapy, regulate haematopoiesis, stimulate wound healing etc. Compositions comprising the polynucleotides may also be administered, especially to express CK beta -13 polypeptide in hosts to treat dysfunctions associated with aberrant endogenous CK beta -13 activity. The polynucleotides are also useful for mapping of chromosomes/chromosome sites. The polypeptides are useful to screen for agonists and antagonists of CK beta -13 activity. The antibodies are useful **diagnostically** or therapeutically e.g. as antagonists to treat subjects requiring CK beta -13 reduction.

Dwg.0/4

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DOC. NO. NON-CPI: N1998-225472
DOC. NO. CPI: C1998-088988
TITLE: Lung tissue derived polynucleotide LU103 - useful
to detect, diagnose, stage,
monitor, prognose, prevent, treat or
determine pre-disposition to lung
disease, e.g. lung cancer.
DERWENT CLASS: B04 D16 S03
INVENTOR(S): COHEN, M; FRIEDMAN, P N; GORDON, J; HODGES, S C;
KLASS, M R; KRATOCHVILL, J D; ROBERTS-RAPP, L;
RUSSELL, J C; STROUPE, S D
PATENT ASSIGNEE(S): (ABBO) ABBOTT LAB
COUNTRY COUNT: 19
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9820143	A1	19980514	(199825)*	EN	86
RW: AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE					
W: CA JP					

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 9820143	A1	WO 1997-US20680	19971105

PRIORITY APPLN. INFO: US 1996-744211 19961105

AN 1998-286957 [25] WPIDS

AB WO 9820143 A UPAB: 19980715

The following are claimed: (1) a method for detecting the presence of a target LU103 polynucleotide in a test sample, comprising: (a) contacting the sample with at least 1 LU103-specific polynucleotide, and (b) detecting the target LU103 polynucleotide in the test sample, where the LU103 polynucleotide has at least 50% identity to the 269, 263, 225, 507 or 519 bp nucleic acid sequence given in the specification; (2) a method for detecting LU103 mRNA in a test sample, comprising: (a) performing reverse transcription with at least 1 primer in order to produce cDNA; (b) amplifying the cDNA using LU103 oligonucleotides as sense and antisense primers to obtain LU103 amplicon, and (c) detecting LU103 amplicon in the test sample, where the LU103 oligonucleotides utilised in steps (a) and (b) have at least 50% sequence identity to the 269, 263, 225, 507 or 519 bp sequence and (3) a method detecting a target LU103 polynucleotide in a test sample suspended of containing the target, comprising: (a) contacting the test sample with at least 1 LU103 oligonucleotide as a sense primer and at least 1 LU103 oligonucleotide as an anti-sense primer and amplifying to obtain a

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first stage reaction product; (b) contacting the first stage reaction product with at least 1 other LU103 oligonucleotide to obtain a second stage reaction product, provided that the other LU103 oligonucleotide is located 3' to the LU103 oligonucleotides utilised in step (a) and is complementary to the first stage reaction product, and (c) detecting the second stage reaction product as an indication of the presence of the target LU103 polynucleotide, where the LU103 oligonucleotides utilised in steps (a) and (b) have at least 50% sequence identity to the 269, 263, 225, 507 or 519 bp sequence; (4) a purified polynucleotide derived from an LU103 gene, where the polynucleotide is capable of selectively **hybridising** to the nucleic acid of the LU103 gene and has at least 50% identity to the 269, 263, 225, 507 or 519 bp sequence; (5) a recombinant expression system comprising a nucleic acid sequence that includes an ORF derived from LU103 operably linked to a control sequence compatible with a desired host, where the nucleic acid sequence has at least 50% identity to the 269, 263, 225, 507 or 519 bp sequence; (6) cell transfected with the recombinant expression system; (7) cell transfected with a nucleic acid sequence encoding at least 1 LU103 **epitope**, where the nucleic acid sequence has the 269, 263, 225, 507 or 519 bp sequence; (8) composition comprising a LU103 polynucleotide, where the polynucleotide has at least 50% identity to the 269, 263, 225, 507 or 519 bp sequence; (9) gene which encodes a LU103 protein which comprises an amino acid sequence with at least 50% identity with the 93 residue amino acid sequence given in the specification, and (10) gene comprising DNA having at least 50% identity with the 507 or 519 bp sequence.

USE - The methods and products of the invention may be used to detect, diagnose, stage, monitor, prognose, prevent, treat or determine the predisposition diseases and conditions of the lung, e.g. lung cancer.

Dwg.0/4

(FILE 'USPATFULL' ENTERED AT 12:18:19 ON 25 OCT 1999)

L9 7 S L2(S) (REAGENT OR EPITOPE OR LS147 OR LS 147)

L9 ANSWER 1 OF 7 USPATFULL

ACCESSION NUMBER: 1999:96216 USPATFULL

TITLE: **Reagents and methods useful for
detecting diseases of the
lung**

INVENTOR(S): Cohen, Maurice, Highland Park, IL, United States
Friedman, Paula N., Deerfield, IL, United States
Gordon, Julian, Lake Bluff, IL, United States
Hodges, Steven C., Buffalo Grove, IL, United States
Klass, Michael R., Libertyville, IL, United States

Searcher : Shears 308-4994

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Kratochvil, Jon D., Kenosha, WI, United States
Roberts-Rapp, Lisa, Gurnee, IL, United States
Russell, John C., Kenosha, WI, United States
Stroupe, Steven D., Libertyville, IL, United States
PATENT ASSIGNEE(S): Abbott Laboratories, Abbott Park, IL, United States (U.S. corporation)

	NUMBER	DATE
PATENT INFORMATION:	US 5939265	19990817
APPLICATION INFO.:	US 1997-964725	19971105 (8)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1996-744211, filed on 5 Nov 1996, now abandoned	
DOCUMENT TYPE:	Utility	
PRIMARY EXAMINER:	Degren, Nancy	
ASSISTANT EXAMINER:	Wang, Andrew	
LEGAL REPRESENTATIVE:	Becker, Cheryl L.; Goller, Mimi C.	
NUMBER OF CLAIMS:	21	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	6 Drawing Figure(s); 6 Drawing Page(s)	
LINE COUNT:	3052	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A set of contiguous and partially overlapping RNA sequences and polypeptides encoded thereby, designated as LU103 and transcribed from lung tissue is described. A fully sequenced clone representing the longest continuous sequence of LU103 is also disclosed. These sequences are useful for detecting, diagnosing, staging, monitoring, prognosticating, preventing or treating, or determining the predisposition of an individual to diseases and conditions of the lung such as lung cancer.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

INCL INCLM: 435/006.000
INCLS: 435/172.300; 435/320.100; 435/325.000; 536/023.100;
536/023.500
NCL NCLM: 435/006.000
NCLS: 435/320.100; 435/325.000; 536/023.100; 536/023.500

L9 ANSWER 2 OF 7 USPATFULL

ACCESSION NUMBER: 1999:43653 USPATFULL
TITLE: Amidinohydrazones as protease inhibitors
INVENTOR(S): Soll, Richard M., Lawrenceville, NJ, United States
Lu, Tianbao, Exton, PA, United States
Fedde, Cynthia L., Warrington, PA, United States
Tomczuk, Bruce E., Collegeville, PA, United States
Illig, Carl, Phoenixville, PA, United States
Searcher : Shears 308-4994

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PATENT ASSIGNEE(S): 3-Dimensional Pharmaceuticals, Inc., Exton, PA,
United States (U.S. corporation)

	NUMBER	DATE
PATENT INFORMATION:	US 5891909	19990406
APPLICATION INFO.:	US 1997-828160	19970327 (8)

	NUMBER	DATE
PRIORITY INFORMATION:	US 1996-14317	19960329 (60)
DOCUMENT TYPE:	Utility	
PRIMARY EXAMINER:	Richter, Johann	
ASSISTANT EXAMINER:	Oswecki, Jane C.	
LEGAL REPRESENTATIVE:	Sterne, Kessler, Goldstein & Fox P.L.L.C.	
NUMBER OF CLAIMS:	76	
EXEMPLARY CLAIM:	1	
LINE COUNT:	5001	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Amidino and benzamidino compounds, including compounds of the formula: ##STR1## wherein R.sup.1 -R.sup.4, R.sup.6 -R.sup.9, Y, Z, n and m are set forth in the specification, as well as hydrates, solvates or pharmaceutically acceptable salts thereof, that inhibit proteolytic enzymes such as thrombin are described. Also described are methods for preparing the compounds of Formula I.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

INCL INCLM: 514/517.000
INCLS: 514/309.000; 514/312.000; 514/345.000; 514/361.000;
514/401.000; 514/406.000; 514/518.000; 514/530.000;
514/567.000; 514/632.000; 546/141.000; 546/172.000;
546/290.000; 548/127.000; 548/353.100; 548/367.100;
549/230.000; 558/056.000; 560/303.000; 562/431.000;
562/432.000; 564/228.000

NCL NCLM: 514/517.000
NCLS: 514/309.000; 514/312.000; 514/345.000; 514/361.000;
514/401.000; 514/406.000; 514/518.000; 514/530.000;
514/567.000; 514/632.000; 546/141.000; 546/172.000;
546/290.000; 548/127.000; 548/353.100; 548/367.100;
549/230.000; 558/056.000; 560/303.000; 562/431.000;
562/432.000; 564/228.000

L9 ANSWER 3 OF 7 USPATFULL

ACCESSION NUMBER: 1998:157475 USPATFULL
TITLE: Hybridomas producing monoclonal antibodies to new
mucin epitopes
INVENTOR(S): Linsley, Peter S., Seattle, WA, United States
Horn, Diane, Seattle, WA, United States
Searcher : Shears 308-4994

09/092296

PATENT ASSIGNEE(S): Brown, Joseph P., Seattle, WA, United States
Sanofi, Paris, France (non-U.S. corporation)

	NUMBER	DATE
PATENT INFORMATION:	US 5849876	19981215
APPLICATION INFO.:	US 1994-179875	19940111 (8)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1992-913740, filed on 14 Jul 1992, now abandoned which is a continuation of Ser. No. US 1987-104511, filed on 8 Oct 1987, now abandoned which is a continuation-in-part of Ser. No. US 1986-932781, filed on 19 Nov 1986, now abandoned	

DOCUMENT TYPE: Utility
PRIMARY EXAMINER: Scheiner, Laurie
LEGAL REPRESENTATIVE: Merchant, Gould, Smith, Edell, Welter & Schmidt
NUMBER OF CLAIMS: 3
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)
LINE COUNT: 1849

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Novel hybridoma cell lines producing monoclonal antibodies reactive with purified mucin antigens from normal and tumor sources are generated using mucins, including purified mucins from tumor sources. Epitopes on mucin antigens from normal and tumor souPPLICATION INFO.: US 1996-698401 19960815 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1995-536939, filed on 29 Sep 1995, now abandoned

DOCUMENT TYPE: Utility
PRIMARY EXAMINER: Ivy, C. Warren
ASSISTANT EXAMINER: Covington, Raymond
LEGAL REPRESENTATIVE: Sterne, Kessler Goldstein & Fox P.L.L.C.
NUMBER OF CLAIMS: 49
EXEMPLARY CLAIM: 1
LINE COUNT: 4363

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Compounds of the formula: ##STR1## wherein R.sup.1 --R.sup.4, R.sup.7 --R.sup.8, R.sup.a, R.sup.b, R.sup.c, Y, Z, n and m are set forth in the specification, as well as hydrates, solvates or pharmaceutically acceptable salts thereof, that inhibit a number of proteolytic enzymes are described. Also described are methods for preparing the compounds of Formula I.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

INCL INCLM: 514/255.000
INCLS: 514/317.000; 514/330.000; 514/331.000; 514/603.000;
514/604.000; 514/634.000; 514/822.000; 544/398.000;
544/399.000; 544/400.000; 544/402.000; 546/021.000;
546/229.000; 546/230.000; 546/231.000; 564/237.000
Searcher : Shears 308-4994

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NCL NCLM: 514/255.000
NCLS: 514/317.000; 514/330.000; 514/331.000; 514/603.000;
514/604.000; 514/634.000; 514/822.000; 544/398.000;
544/399.000; 544/400.000; 544/402.000; 546/021.000;
546/229.000; 546/230.000; 546/231.000; 564/237.000

L9 ANSWER 5 OF 7 USPATFULL

ACCESSION NUMBER: 1998:69171 USPATFULL
TITLE: Cell cycle regulatory gene
INVENTOR(S): Sidransky, David, Baltimore, MD, United States
PATENT ASSIGNEE(S): The Johns Hopkins University School of Medicine,
Baltimore, MD, United States (U.S. corporation)

	NUMBER	DATE
PATENT INFORMATION:	US 5767258	19980616
APPLICATION INFO.:	US 1995-439962	19950512 (8)
DOCUMENT TYPE:	Utility	
PRIMARY EXAMINER:	Elliot, George C.	
ASSISTANT EXAMINER:	McGarry, Sean	
LEGAL REPRESENTATIVE:	Fish & Richardson, P.C.	
NUMBER OF CLAIMS:	9	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	5 Drawing Figure(s); 3 Drawing Page(s)	
LINE COUNT:	1506	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A novel cell cycle regulatory gene called 5'ALT is disclosed.
Methods for determining mutations or polymorphisms in 5'ALT or
5'ALT regulated genes in tissues are also provided. Novel
5'ALT-p16 and 5'ALT-p15 transcripts and truncated expression
products are also described.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

INCL INCLM: 536/023.100
INCLS: 536/023.500; 435/006.000; 435/172.100; 435/320.100;
435/325.000; 435/243.000
NCL NCLM: 536/023.100
NCLS: 435/006.000; 435/243.000; 435/320.100; 435/325.000;
536/023.500

L9 ANSWER 6 OF 7 USPATFULL

ACCESSION NUMBER: 85:8964 USPATFULL
TITLE: .alpha.-Aminoboronic acid peptides
INVENTOR(S): Shenvi, Ashokkumar B., Wilmington, DE, United
States
Kettner, Charles A., Wilmington, DE, United
States
PATENT ASSIGNEE(S): E. I. Du Pont de Nemours and Company, Wilmington,
DE, United States (U.S. corporation)
Searcher : Shears 308-4994

09/092296

	NUMBER	DATE
PATENT INFORMATION:	US 4499082	19850212
APPLICATION INFO.:	US 1983-558362	19831205 (6)
DOCUMENT TYPE:	Utility	
PRIMARY EXAMINER:	Phillips, Delbert R.	
LEGAL REPRESENTATIVE:	Hallquist, Scott G.	
NUMBER OF CLAIMS:	19	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	3 Drawing Figure(s); 3 Drawing Page(s)	
LINE COUNT:	1528	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Peptides comprising C-terminal .alpha.-aminoboronic acid residues are potent, reversible inhibitors of proteolytic enzymes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

INCL INCLM: 514/002.000
INCLS: 260/112.500R; 514/018.000; 514/019.000
NCL NCLM: 514/002.000
NCLS: 260/001.000; 514/018.000; 514/019.000; 514/506.000;
530/331.000; 562/007.000; 930/010.000; 930/023.000;
930/250.000

L9 ANSWER 7 OF 7 USPATFULL

ACCESSION NUMBER: 85:3359 USPATFULL
TITLE: Determination of oxidized .alpha.-1-proteinase inhibitor in serum or plasma
INVENTOR(S): Travis, James, Athens, GA, United States
PATENT ASSIGNEE(S): University of Georgia Research Foundation, Inc., Athens, GA, United States (U.S. corporation)

	NUMBER	DATE
PATENT INFORMATION:	US 4493891	19850115
APPLICATION INFO.:	US 1982-402442	19820727 (6)
DOCUMENT TYPE:	Utility	
PRIMARY EXAMINER:	Kepplinger, Esther M.	
LEGAL REPRESENTATIVE:	Oblon, Fisher, Spivak, McClelland & Maier	
NUMBER OF CLAIMS:	13	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 1 Drawing Page(s)	
LINE COUNT:	603	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A new method of determining oxidized .alpha.-1-proteinase inhibitor in serum or plasma for use in studying the development of chronic obstructive lung disease is disclosed. Levels of oxidized .alpha.-1-proteinase inhibitor indicate the potential for emphysema development with higher levels showing a decrease in

Searcher : Shears 308-4994



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lung protection against elastolytic enzymes such as elastase. This method can be used for patients with a potential for chronic obstructive lung disease rather than having to use bronchial lavage methods for such patients. No other method is known to exist for determining oxidized .alpha.-1-proteinase inhibitor in serum or plasma.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

INCL INCLM: 435/023.000

INCLS: 435/184.000

NCL NCLM: 435/023.000

NCLS: 435/184.000

(FILE 'CAPLUS, MEDLINE, BIOSIS, EMBASE, LIFESCI, WPIDS, CONFSCI, SCISEARCH, JICST-EPLUS, PROMT, CANCERLIT, USPATFULL' ENTERED AT 12:20:18 ON 25 OCT 1999)

L10 134 SEA ABB=ON PLU=ON (BILLING MEDEL P? OR BILLING P? OR
MEDEL P? OR MEDEL BILLING P?)/AU
L11 26222 SEA ABB=ON PLU=ON COHEN M?/AU
L12 123 SEA ABB=ON PLU=ON COLPITTS T?/AU
L13 2767 SEA ABB=ON PLU=ON FRIEDMAN P?/AU
L14 262 SEA ABB=ON PLU=ON KLASS M?/AU
L15 12902 SEA ABB=ON PLU=ON RUSSELL J?/AU
L16 310 SEA ABB=ON PLU=ON STROUPE S?/AU
L17 48 SEA ABB=ON PLU=ON L10 AND L11 AND L12 AND L13 AND L14
AND L15 AND L16
L18 51 SEA ABB=ON PLU=ON L10 AND (L11 OR L12 OR L13 OR L14 OR
L15 OR L16)
L19 62 SEA ABB=ON PLU=ON L11 AND (L12 OR L13 OR L14 OR L15)
L20 51 SEA ABB=ON PLU=ON L12 AND (L13 OR L14 OR L15)
L21 62 SEA ABB=ON PLU=ON L13 AND (L14 OR L15)
L22 59 SEA ABB=ON PLU=ON L14 AND L15

L26 7 SEA ABB=ON PLU=ON (L17 OR L18 OR L19 OR L20 OR L21 OR
L22) AND L1
L27 4 DUP REM L26 (3 DUPLICATES REMOVED)

44

Oct 25 11:54:02 1999

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Tel: 3018699056
Fax: 3018699423
Email: arkerlav@tigr.orgFor clone availability, additional sequence and expression
information related to this EST, please check the TIGR Human Gene
Index (<http://www.tigr.org/tdb/hgi/hgi.html>)
Seq Primer: M13 Reverse.

FEATURES

Location/Qualifiers

1. 311

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ORIGIN

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RESULT

8

LOCUS

AA323964 311 bp mRNA

DEFINITION

EST6816 Cerebellum II Homo sapiens CDNA 5' end, mRNA sequence.

ACCESSION

AA323964

NID

G1976290

VERSION

AA323964.1 GI:1976290

KEYWORDS

EST.

SOURCE

human.

ORGANISM

Homo sapiens

REFERENCE

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

AUTHORS

Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE

1 (bases 1 to 311)

AUTHORS

Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,A.A.,

REFERENCE

Bult,C.J., Lee,N.H., Kirkness,E.F., Weissstock,K.G., Gocayne,J.D.,

AUTHORS

White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-Val,C.,

REFERENCE

Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,

AUTHORS

Fitzgerald,L.M., Fitzgerald,M.M., Fritchman,J.L., Georghiou,N.S.,

REFERENCE

Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S., Jr.,

AUTHORS

Kelley,J.M., Kelley,J.C., Liu,L.-I., Marmaros,S.N., Merrick,J.M.,

REFERENCE

Moreno-Palancas,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M.,

AUTHORS

Phillips,C.A., Ryder,S.E., Scott,J.L., Sauder,D.M., Shirley,R.,

REFERENCE

Small,K.V., Spriggs,T.A., Utterback,T.R., Weidman,J.F., Li,Y.,

AUTHORS

Bednarek,D.P., Cao,D., Cepeda,M.A., Coleman,T.A., Collins,E.J.,

REFERENCE

Dinke,D., Feng,D.-F., Ferris,A., Fischer,C., Hasling,G.A.,

AUTHORS

He,M.W., Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K.,

REFERENCE

Kozak,D.L., Kunsch,C., Hungjun,J., Li,H., Melsner,P.S., Olsen,H.,

AUTHORS

Raymond,L., Wei,Y.F., Wang,J., Xu,C., Yu,G.L., Ruben,S.M.,

REFERENCE

Dillon,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C.,

AUTHORS

Fraser,C.M. and Venter,J.C.

REFERENCE

Initial assessment of human gene diversity and expression patterns

AUTHORS

based upon 83 million nucleotides of cDNA sequence

REFERENCE

Nature 377 (6547 suppl), 3-174 (1995)

AUTHORS

96026280

REFERENCE

On Apr 14, 1993 this sequence version replaced gi:693635.

AUTHORS

Contact: Kerlavage, AR

REFERENCE

Bioinformatics

AUTHORS

The Institute for Genomic Research

REFERENCE

9712 Medical Center Drive, Rockville, MD 20850 USA


```

RESULT 10
LOCUS AA323964 311 bp mRNA EST 20-APR-1997
DEFINITION EST26816 Cerebellum II Homo sapiens cDNA 5' end, mRNA sequence.
ACCESSION AA323964
NID g1976290
VERSION AA323964.1 GI:1976290
KEYWORDS EST.
SOURCE human.
  ORGANISM Homo sapiens
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
    Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 311)
AUTHORS Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A.,
  Bult,C.J., Lee,N.H., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D.,
  White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-Wai,C.,
  Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,
  Fitzgerald,L.M., Fitzhugh,W.M., Fritchman,J.L., Geoghagen,N.S.,
  Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S.Jr.,
  Kelley,J.M., Kelley,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M.,
  Moreno-Palanques,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M.,
  Phillips,C.A., Ryder,S.E., Scott,J.L., Saudek,D.M., Shirley,R.,
  Small,K.V., Spriggs,T.A., Utterback,T.R., Weidman,J.F., Li,Y.,
  Bednarik,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,
  Dimke,D., Feng,D.-F., Ferrie,A., Fischer,C., Hastings,G.A.,
  He,W.W., Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K.,
  Kozak,D.L., Kunsch,C., Hungjun,J., Li,H., Meissner,P.S., Olsen,H.,
  Raymond,L., Wei,Y.F., Wing,J., Xu,C., Yu,G.L., Ruben,S.M.,
  Dillion,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C.,
  Fraser,C.M. and Venter,J.C.
  TITLE Initial assessment of human gene diversity and expression patterns
  JOURNAL Nature 377 (6547 Suppl), 3-174 (1995)
  MEDLINE 96026280
  COMMENT On Apr 14, 1993 this sequence version replaced gi:693635.
    *
    Contact: Kerlavage, AR
    Bioinformatics
    The Institute for Genomic Research
    9712 Medical Center Drive, Rockville, MD 20850 USA
    Tel: 3018699056
    Fax: 3018699423
    Email: arkerlav@tigr.org
    For clone availability, additional sequence and expression
    information related to this EST, please check the TIGR Human Gene
    Index (http://www.tigr.org/tdb/hgi/hgi.html)
    Seq primer: M13 Reverse.
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    ||||| |||| |||||
  Qy 43 TGGCCACTATGGGGTCTGGGCTGCCCTT 71

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Oct 25 11:53:59 1999

US-09-092-296-3.mri

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TELEX: 24855 OPAT UR
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 3088 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
SEQUENCE 3088 BP: 716 A; 761 C; 672 G; 939 T; 0 OTHER.

Query Match 12.24; Score 22; DB 3; Length 3088;
Best Local Similarity 72.04; Pred. No. 7,70e-02;
Matches 36; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

15 CCGAGATGGGCCAGAGGGGGCGCTGCCCGAGGCGCTGCTGCGC 64
63 CCAAGAGGCTCAAGAGGAGGACAGGCGACCCCATAGTGGC 14

RESULT 8
ID US-08-418-444A-1 STANDARD; ONA; UNC; 3088 BP.
AC xxxxxx
DT Sequence 1, Application US/08418444A
CC Sequence 1, Application US/08418444A
CC Patent No. 5773688
CC GENERAL INFORMATION:
CC APPLICANT: KURODA, HISAO
CC APPLICANT: RIKOTA, NAORIKO
CC TITLE OF INVENTION: GENE EXPRESSION VECTOR USING THE GENE
CC TITLE OF INVENTION: GENE EXPRESSION REGULATING REGION OF THE ADP RIBOSYLATION
CC TITLE OF INVENTION: FACTOR
CC NUMBER OF SEQUENCES: 9
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: OHLON, SPYVAK, MCCLELLAND, MAIER & NEUSTADT
CC STREET: 1755 S. JEFFERSON DAVIS HIGHWAY, SUITE 400
CC CITY: ARLINGTON
CC STATE: VIRGINIA
CC COUNTRY: USA
CC ZIP: 22202
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patent Release #1.0, Version #1.30
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/418,444A
CC FILING DATE: 07-APR-1995
CC CLASSIFICATION: 800
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: JP HEI 6-71048
CC FILING DATE: 08-APR-1994
CC ATTORNEY/AGENT INFORMATION:
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CC REFERENCE/DOCKET NUMBER: 2589-024-0
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